

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 30, 2002, 02:35:01 : Search time 58.67 seconds
(without alignments)

8658.089 Million cell updates/sec

Title: US-09-932-678-1

Perfect score: 2068

Sequence: 1 acgaagacacagcagcaggaag.....tgacatcttgaatccatc 2068

Scoring table: IDENTITY-MNC

Searched: 38533 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 767066

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents NA:*

pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match length	ID	Description
1	48.2	2.3	8797	2	US-08-723-306-6
2	48.2	2.3	8797	5	PCT-US96-10041-6
3	48.2	2.3	11093	2	US-08-723-306-5
4	48.2	2.3	11093	5	PCT-US96-10041-5
5	40.4	2.0	7218	1	US-08-232-464-14
6	39.2	1.9	4937	1	US-08-038-662-3
7	39.2	1.9	4937	1	US-08-302-832-3
8	39.2	1.9	4937	2	US-08-510-158-3
9	39.2	1.9	4937	2	US-08-469-880-3
10	39.2	1.9	4937	2	US-08-728-470-3
11	39.2	1.9	4937	2	US-08-617-697-3
12	39.2	1.9	4937	4	US-08-719-641-3
13	35.2	1.7	5538	2	US-08-231-193A-55
14	35.2	1.7	5538	2	US-08-486-273A-55
15	35.2	1.7	5538	3	US-08-940-086A-55
16	35.2	1.7	5538	4	US-08-940-035A-55
17	34.6	1.7	2518	4	US-09-433-699-3
18	34.4	1.7	289	4	US-09-007-005-17
19	34.4	1.7	289	4	US-09-244-796-17
20	34.4	1.7	1307	4	US-09-282-305-17
21	34.4	1.7	9323	1	US-08-038-682-6
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28	34	1.6	1245	1	US-08-385-186-14	Sequence 14, Appl
C 29	33.4	1.6	10614	1	US-08-135-511-35	Sequence 45, Appl
C 30	33.4	1.6	10614	1	US-08-187-453-35	Sequence 45, Appl
31	33.2	1.6	1545	1	US-07-872-673B-2	Sequence 2, Appl
C 32	33.2	1.6	2454	1	US-07-872-673B-2	Sequence 1, Appl
C 33	32.6	1.6	19182	2	US-08-850-880-11	Sequence 11, Appl
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C 35	32.4	1.6	2791	4	US-09-570-367-1	Sequence 1, Appl
36	32.2	1.6	4856	4	US-09-045-460-1	Sequence 1, Appl
37	32.2	1.6	12225	2	US-08-822-445-11	Sequence 11, Appl
38	32.2	1.6	12225	4	US-09-396-540-11	Sequence 11, Appl
39	32.2	1.6	12616	2	US-08-822-445-9	Sequence 9, Appl
40	32.2	1.6	12616	4	US-09-396-540-9	Sequence 9, Appl
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42	32	1.5	1202	3	US-09-058-489-29	Sequence 29, Appl
C 43	32	1.5	1248	3	US-08-910-505-4	Sequence 4, Appl
C 44	32	1.5	1251	3	US-08-910-505-1	Sequence 1, Appl
C 45	31.8	1.5	1338	3	US-09-027-064-4	Sequence 4, Appl

ALIGNMENTS

RESULT 1
US-08-723-306-6
Sequence 6, Application US/08723306
Patent No. 5856178
GENERAL INFORMATION:
APPLICANT: White PhD, Kenneth
APPLICANT: Morrey PhD, John
APPLICANT: Reed, William
TITLE OF INVENTION: Cassette for expression of lytic
NUMBER OF SEQUENCES: 32
REFERENCE SEQUENCES: 32
ADDRESS: Task Britl and Kossa
STREET: P.O. Box 2550
CITY: Salt Lake City
STATE: Utah
COUNTRY: USA
ZIP: 84110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patcon in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/723,306
FILING DATE:
CLASSIFICATION: 800
ATTORNEY/AGENT INFORMATION:
NAME: Sweigert PhD, Susan E
REGISTRATION NUMBER: 36,289
REFERENCE/DOCKET NUMBER: 2549
TELECOMMUNICATION INFORMATION:
TELEPHONE: 8015321922
TELEFAX: 8015319168
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 8797 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
POPLOGY: not relevant
MOL-VEH TYPE: other nucleic acid
DESCRIPTION: /desc - "Construct comprising fos
DESCRIPTION: tauus beta casein 5' regulatory region plus genes encoding
DESCRIPTION: amphipathic peptide and green fluorescent protein"
HYDRETICAL: YES
ANT-SENSE: NO
US-08-723-306-6
Query Match 2.3%, Score 48.2, DB 2, Length 8797:

[illegible]

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1 RESULT 2
2 PCT 0036 10041 6
3 SEQUENCE 6, Application 167/US9610041
4 GENERAL INFORMATION:
5 APPLICANT: White PhD, Kenneth
6 APPLICANT: Morley PhD, John
7 APPLICANT: Reed, William
8 TITLE OF INVENTION: Cloned for Expression of Lysic
9 TITLE OF INVENTION: Peptides in Mammalian Transgenic Organisms
10 NUMBER OF SEQUENCES: 42
11 CORRESPONDENCE ADDRESS:
12 ADDRESSEE: Frank Hill and Rossa
13 STREET: P.O. Box 2550
14 CITY: Salt Lake City
15 STATE: Utah
16 COUNTRY: USA
17 ZIP: 84110
18 COMPUTER READABLE FORM:
19 MEDIUM TYPE: floppy disk
20 COMPUTER: IBM PC compatible
21 OPERATING SYSTEM: PC DOS/MS DOS
22 SOFTWARE: Patent in Release #1.0, Version #1.40
23 CURRENT APPLICATION DATA:
24 APPLICATION NUMBER: PCT/US96/10041
25 FILING DATE:
26 CLASSIFICATION:
27 AUTOREY/AGENT INFORMATION:
28 NAME: Scottport, Fred, Susan E
29 REGISTRATION NUMBER: 60,289
30 REFERENCE/BOOK NUMBER: 2949
31 TELECOMMUNICATION INFORMATION:
32 TELEPHONE: 8015421922
33 TELEFAX: 8015419168
34 INFORMATION FOR SEQ ID NO: 6:
35 SEQUENCE CHARACTERISTICS:
36 LENGTH: 8797 base pairs
37 TYPE: nucleic acid
38 STANDARDS: GenBank
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41 DESCRIPTION: /desc "construct encoding Bos
42 DESCRIPTION: Laurus beta casein 5' regulatory region plus genes encoding
43 DESCRIPTION: amphipathic peptide and green fluorescent protein"
44 HYDROPHATIC: YES
45 ANTI SENSE: NO
46 PCT 0036 10041 6

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Host Team's Difficulty	52-28	Prod. No.	0-00043			
Matches 197	Consecutive	0	Misses	90	Indels	0
697	Amount of Total	Amount of	Amount of	Amount of	Amount of	Amount of
527	697	Amount of	Amount of	Amount of	Amount of	Amount of

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1 RESIDU 4
2 US-08 754-306 5
3 Sequence 5, Application US/08/724,006
4 Patent No. 5864178
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6 GENERAL INFORMATION:
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8 APPLICANT: White PhD, Kenneth
9 APPLICANT: Morley PhD, John
10 APPLICANT: Reed, William
11 TITLE OF INVENTION: Cassette for Expression of Lytic
12 TITLE OF INVENTION: Peptides in Mammalian Transgenic Organisms
13
14 NUMBER OF SEQUENCES: 42
15 CORRESPONDENCE ADDRESS:
16 ADDRESSEE: Task 1111 and Rossa
17 STREET: P.O. Box 2550
18 CITY: Salt Lake City
19 STATE: Utah
20 COUNTRY: USA
21
22 ZIP: 84110
23
24 COPY, PER READABLE FORM:
25 MEDIUM TYPE: floppy disk
26 COMPUTER: IBM pc compatible
27 OPERATING SYSTEM: pc dos/ms dos
28 SOFTWARE: Patent In Release #1.0, Version #1.40
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30 CURRENT APPLICATION DATA:
31 APPLICATION NUMBER: US/08/724,006
32 FILING DATE:
33 CLASSIFICATION: 800
34 ATTORNEY/AGENT INFORMATION:
35 NAME: Sweett 100, Susan E
36 REGISTRATION NUMBER: 46,289
37 REFERENCE/BOOKLET NUMBER: 2,049
38 TELECOMMUNICATION INFORMATION:
39 TELEPHONE: 8015421922
40 TELEFAX: 8015419168
41 INFORMATION FOR SEQ ID NO: 5:
42 SEQUENCE CHARACTERISTICS:
43 LENGTH: 11094 base pairs
44 TYPE: nucleic acid
45 STRANDEDNESS: double
46 TOPOLOGY: not relevant
47 MOLECULE TYPE: other nucleic acid
48 DESCRIPTION: /desc "Construct comprising
49 DESCRIPTION: portions of boe taurin beta casein gene and genes encoding
50 DESCRIPTION: amphipathic peptide and green fluorescent protein."
51 HYDROPHOBIC: YES
52 ANTI-SENSE: NO
53 FEATURE:
54 NAME/KEY: exon
55 LOCATION: 1801..1834
56 OTHER INFORMATION: /product "beta casein exon 1"
57 FEATURE:
58 NAME/KEY: exon
59 LOCATION: 4780..4832
60 OTHER INFORMATION: /product "boe casein exon 2"
61 FEATURE:
62 NAME/KEY: DATA.signal
63 LOCATION: 1766..1774

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NAME/KEY: exon
LOCATION: 4567..4590
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OTHER INFORMATION: exon 3"
FEATURE:
NAME/KEY: 5'UTR
LOCATION: 127..1800
OTHER INFORMATION: /function= "5' flanking regulatory"
OTHER INFORMATION: region of bovine beta casein gene"
FEATURE:
NAME/KEY: mat_peptide
LOCATION: 4587..5310
OTHER INFORMATION: /product= "Green fluorescent"
OTHER INFORMATION: protein"
FEATURE:
NAME/KEY: mat_peptide
LOCATION: 5320..5449
OTHER INFORMATION: /product= "Shiva-1 coding sequence"
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NAME/KEY: polyA-signal
LOCATION: 7640..7635
OTHER INFORMATION: /standard_name= "Bovine beta casein"
OTHER INFORMATION: 3' region, in exon 9"

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Matches 107	Conservative 0	Mismatches 98	Indels 0	Gaps 0

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Db	2278 AAAAACAAGCATATATTATTTAAAAATTCGTACAGACATCCAAATTTTCAGTTATCTTAT	2333
QY	757 taagatttagtataattttcccaactttagagatgaattctgaagcttaattatga	816
Db	2338 CTTCACAAATATTCGAAAATAATTAATAATAGATACATGAATACGAAATTAATAAGAGA	2397
QY	817 aactactcaagtttgaatgtgaatgcaatcccaagaggtattgaatgtctgaagaacag	876
Db	2398 AAGTATTTTATTTGTAAAAAAAATTTAGTGTGCACGAGGAGTACAGCAAAACAAAA	2457
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Db	2458 ACAGTGAANAATGTGATCTGACGCA 2482	

RESULT 4
 PCT-US96-10041-5
 Sequence 5, Application PC/US9610041
 GENERAL INFORMATION:
 APPLICANT: White PhD, Kenneth
 APPLICANT: Morrey PhD, John
 APPLICANT: Reed, William
 TITLE OF INVENTION: Cassette for Expression of Lytic
 TITLE OF INVENTION: Peptides in Mammalian Transgenic Organisms
 NUMBER OF SEQUENCES: 32
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Trask Britt and Rossa
 STREET: P.O. Box 2550
 CITY: Salt Lake City
 STATE: Utah
 COUNTRY: USA
 ZIP: 84110
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patent In Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: PCT/US96/10041
 FILING DATE:

	CLASSIFICATION:
	ATTORNEY/AGENT INFORMATION:
	NAME: Sweigert Ph.D. Susan E
	REGISTRATION NUMBER: 46,289
	REFERENCE/DOCKET NUMBER: 2549
	TELECOMMUNICATION INFORMATION:
	TELEPHONE: 8015321922
	TELEFAX: 8015319168
	INFORMATION FOR SEQ ID NO: 5:
	SEQUENCE CHARACTERISTICS:
	LENGTH: 1109 base pairs
	TYPE: nucleic acid
	STRANDEDNESS: double
	TOPOLOGY: not relevant
	MOLECULE TYPE: other nucleic acid
	DESCRIPTION /desc = "construct comprising
	DESCRIPTION: portions of Bos taurus beta casein gene and sheep cytochrome
	DESCRIPTION: amphipathic peptide and green fluorescent protein"
	HYPOTHETICAL: YES
	ANTI-SENSE: NO
	FEATURE:
	NAME/KEY: exon
	LOCATION: 1801..1844
	OTHER INFORMATION: /product "beta casein exon 1"
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	OTHER INFORMATION: /product - "beta casein exon 2"
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	NAME/KEY: TATA_signal
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	NAME/KEY: exon
	LOCATION: 4567..4590
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	OTHER INFORMATION: exon 3"
	FEATURE:
	NAME/KEY: 5'UTR
	LOCATION: 127..1800
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	OTHER INFORMATION: region of bovine beta casein gene"
	FEATURE:
	NAME/KEY: mat_peptide
	LOCATION: 4587..5310
	OTHER INFORMATION: /product - "green fluorescent
	OTHER INFORMATION: protein"
	FEATURE:
	NAME/KEY: mat_peptide
	LOCATION: 5320..5449
	OTHER INFORMATION: /product - "Shiva-1 coding sequence"
	FEATURE:
	NAME/KEY: polyA_signal
	LOCATION: 7640..7635
	OTHER INFORMATION: /standard_name "bovine beta casein
	OTHER INFORMATION: 3' region, in exon 9"
	PCT-US95-19041-5
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Best local Similarity	52.2%; Iter: No; 0,00047;
Matches 107; Conservative 0; Mismatches 98; Indels 0; Gaps 0;	
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Dy 2276	AAAGGAGGATATTTTATTAAATTCGTACAGACTGCMAATTTTAATTTAT 2397
Dy 757	taaqatatatataatttcgaacctgaagcacaatctgaqatctaatat 816
Dy 2338	CTTGCAATATTTCCAAAAATATTTAAATATGATCATGAATATACAAATTAAT 2397
Dy 817	aagtaactcaagttagatgtgatgtatgcccgacgaagatdtaaaatgctaaaataatg 876
Dy 2395	AAGTATTTTATTTTCTAAAAAAAATTCCTACCTGGCATAGGCAATACAGCAAA 2457


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1  STREET: 2001 JOHNSON DAVIS HWY., 1204 CRYSTAL PLAZA
2  CITY: ATLANTION
3  STATE: VIRGINIA
4  COUNTRY: U.S.A.
5  ZIP: 2202 0286
6  COMPUTER READABLE FORM:
7  MEDIUM TYPE: floppy disk
8  COMPUTER: IBM pc compatible
9  OPERATING SYSTEM: pc dos/MS DOS
10 SOFTWARE: Paton In Release #1.0, Version #1.00
11 CURRENT APPLICATION DATA:
12 APPLICATION NUMBER: HS/08/469,080
13 FILING DATE: 06 JUN-1995
14 CLASSIFICATION: 435
15 PRIOR APPLICATION DATA:
16 APPLICATION NUMBER: GB 9205704.1
17 FILING DATE: 16 MAR 1992
18 PRIOR APPLICATION DATA:
19 APPLICATION NUMBER: US PAT/HS/402166
20 FILING DATE: 16 MAR 1993
21 PRIOR APPLICATION DATA:
22 APPLICATION NUMBER: US 08/402,832
23 FILING DATE: 16 SEP 1994
24 ATTORNEY/AGENT INFORMATION:
25 NAME: BOKSTONSON, JERRY W
26 REGISTRATION NUMBER: 22,651
27 REFERENCE/PACKET NUMBER: 1048 516 MIS:V01
28 TELECOMMUNICATION INFORMATION:
29 TELEPHONE: (703) 415,0810
30 TELEFAX: (703) 415,0814
31 INFORMATION FOR SEQ ID NO: 4:
32 SEQUENCE CHARACTERISTICS:
33 LENGTH: 4997 base pairs
34 TYPE: nucleic acid
35 STRANDEDNESS: single
36 TOPOLOGY: linear
37 MOLECULE TYPE: DNA (genomic)
38 DS DB 469 860 4

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COUNTRY: U.S.A.
 ZIP: 22202-0286
 COMPUTER READABLE FORM:
 MEDIUM TYPE: floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patent in Release #1.0, Version #1.40
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/728,470
 FILING DATE:
 CLASSIFICATION: 424
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/402,842
 FILING DATE: 16-MAR-1994
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US PCT/0834/02166
 FILING DATE: 16-MAR-1994
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: GR 9205704.1
 FILING DATE: 16-MAR-1992
 ATTORNEY/AGENT INFORMATION:
 NAME: HOKSTADT, JEFF W
 REGISTRATION NUMBER: 22,661
 REFERENCE/DOCKET NUMBER: 1038-643
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (703) 415-0810
 TELEFAX: (703) 415-0814
 INFORMATION FOR SEQ ID NO: 4:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 4997 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 topology: linear
 MOLECULE TYPE: DNA (genomic)
 OS-08-728-470-4

[illegible]

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1      PRIOR APPLICATION DATA:
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6      APPLICATION NUMBER:  US PAT/0894/02166
7      FILING DATE:  16-MAR-1993
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9      PRIOR APPLICATION DATA:
10     APPLICATION NUMBER:  GR 9205704.1
11     FILING DATE:  16-MAR-1992
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13     ATTORNEY/AGENT INFORMATION:
14     NAME:  Borkstrosser, Jerry W
15     REGISTRATION NUMBER:  22,651
16     REFERENCE/DOCKET NUMBER:  1038-625
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18     TELECOMMUNICATION INFORMATION:
19     TELEPHONE:  (703) 415-0810
20     TELEFAX:  (703) 415-0813
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OY	803	tccatattgaaactacacgaattgataagaaagcctcaagaatttgaad	862
Db	4287	tggttttaaaccaaaagtcttagcttaaatgcctatgatcatgccgc	4346
OY	863	tgttgaagaataggcacctcaacctgttgtggagaaatttcacagaaattgttaa	922
Db	4447	saatgcacattaacaccanagcgctcttgatgtatgcattgcgggaaccttcanaagattttaa	4406
OY	923	tatgatgaagatgaauaaattgaacatgaacaacaa	958
Db	4467	tattactgcacatatlaaacatcatttaaatgcggttaa	4442

RESULT 14
 US-08-231-193A-55
 Sequence 55, Application US/08231193A
 Patent No. 5849895
 GENERAL INFORMATION:
 APPLICANT: Daggett, Lottie P.
 APPLICANT: Ellis, Steven B.
 APPLICANT: Liaw, Chen W.
 APPLICANT: Lu, Chin-Chou
 TITLE OF INVENTION: HUMAN N-METHYL-D-ASPARTATE RECEPTOR
 TITLE OF INVENTION: SUBUNITS, NOCTEIC ACIDS EMBODING SAME AND USES THEREOF
 NUMBER OF SEQUENCES: 63
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Brown, Martin, Hallor & McElain
 STREET: 1660 Union Street
 CITY: San Diego
 STATE: CA
 COUNTRY: U.S.A.
 ZIP: 92101-2926
 COMPUTER READABLE FORM:
 MEDIUM TYPE: floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patent in release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/231,193A
 FILING DATE: 20-APR-1994
 CLASSIFICATION: 536
 PUBLICATION DATA:
 PUBLICATION NUMBER: US 08/052,459

1 FILING DATE: 20 APR 1994
 2 CLASSIFICATION: 546
 3 AT/PATENT/AGENT INFORMATION:
 4 NAME: Solomon, Stephanie
 5 REGISTRATION NUMBER: 44,779
 6 REFERENCE/PATENT NUMBER: 6,662,948
 7 TELEPHONE: 619-238-0999
 8 TELEFAX: 619-238-0062
 9 INFORMATION FOR SEQ ID NO: 55:
 10 SEQUENCE CHARACTERISTICS:
 11 LENGTH: 5548 base pairs
 12 TYPE: nucleic acid
 13 STRANDEDNESS: both
 14 TOPOLOGY: both
 15 FEATURE:
 16 NAME/KEY: CDS
 17 LOCATION: 210..4664
 18 ORF 486 274A 55

Query Match 1.7% Score 45.2; DB 2; Length 5548;
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 Matches 88; Conservative 0; Mismatches 88; Indels 0; Gaps 0;

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 QY 1822 atgtctgaagagctcaagaaatctatgactatctatcagatctgaagagagca 1881
 DB 1433 GATTTCATCTGTCGAAAGCTGAGAGCTGAGATCTGAGATCTGAGATCTGAGATCT 1452
 QY 1882 atgtctgaagagctcaagaaatctatgactatctatcagatctgaagagagca 1937
 DB 1433 GATTTCATCTGTCGAAAGCTGAGAGCTGAGATCTGAGATCTGAGATCTGAGATCT 1508

1 RESULT 14
 2 US 08 486 274A 55
 3 Sequence 55, Application US/0848627A
 4 Patent No. 6,662,948
 5 GENERAL INFORMATION:
 6 APPLICANT: Baggott, Louie P.
 7 APPLICANT: Ellis, Steven B.
 8 APPLICANT: Liaw, Chen W.
 9 APPLICANT: Lu, Chiu-Chun
 10 TITLE OF INVENTION: HUMAN N-METHYL-D-ASPARTATE RECEPTOR SUBUNITS, DNA
 11 TITLE OF INVENTION: ENCODING SAME AND USES THEREOF
 12 NUMBER OF SEQUENCES: 64
 13 ADDRESS/INVENTOR ADDRESS:
 14 ADDRESSEE: Brown, Martin; Bullock & McCallan
 15 STREET: 1660 Union Street
 16 CITY: San Diego
 17 STATE: CA
 18 COUNTRY: U.S.A.
 19 ZIP: 92101 2926
 20 COMPUTER READABLE FORM:
 21 MEDIUM TYPE: Floppy disk
 22 COMPUTER: IBM PC compatible
 23 OPERATING SYSTEM: PC-DOS/MS-DOS
 24 SOFTWARE: Patent In Release #1.0, Version #1.25
 25 CURRENT APPLICATION DATA:
 26 APPLICATION NUMBER: US/08/486,27A
 27 FILING DATE: 06-JUN-1995
 28 CLASSIFICATION: 435
 29 PRIOR APPLICATION DATA:
 30 APPLICATION NUMBER: US 08/231,193
 31 FILING DATE: 20 APR 1994
 32 CLASSIFICATION: 435
 33 ATTORNEY/AGENT INFORMATION:
 34 NAME: Solomon, Stephanie

1 REGISTRATION NUMBER: 44,779
 2 REFERENCE/PATENT NUMBER: 6,662,948
 3 TELECOMMUNICATION INFORMATION:
 4 TELEPHONE: 619-238-0999
 5 TELEFAX: 619-238-0062
 6 INFORMATION FOR SEQ ID NO: 55:
 7 SEQUENCE CHARACTERISTICS:
 8 LENGTH: 5548 base pairs
 9 TYPE: nucleic acid
 10 STRANDEDNESS: both
 11 TOPOLOGY: both
 12 FEATURE:
 13 NAME/KEY: CDS
 14 LOCATION: 210..4664
 15 ORF 486 274A 55

Query Match 1.7% Score 45.2; DB 2; Length 5548;
 Best Local Similarity 50.0%; Pred. No. 2.5;
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QY 1762 atgtctgaagagctcaagaaatctatgactatctatcagatctgaagagagca 1821
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 QY 1822 atgtctgaagagctcaagaaatctatgactatctatcagatctgaagagagca 1881
 DB 1433 GATTTCATCTGTCGAAAGCTGAGAGCTGAGATCTGAGATCTGAGATCTGAGATCT 1452
 QY 1882 atgtctgaagagctcaagaaatctatgactatctatcagatctgaagagagca 1937
 DB 1433 GATTTCATCTGTCGAAAGCTGAGAGCTGAGATCTGAGATCTGAGATCTGAGATCT 1508

1 RESULT 15
 2 US 08 940 086A 55
 3 Sequence 55, Application US/08940086A
 4 Patent No. 6,111,091
 5 GENERAL INFORMATION:
 6 APPLICANT: Baggott, Louie P.
 7 APPLICANT: Ellis, Steven B.
 8 APPLICANT: Liaw, Chen W.
 9 APPLICANT: Lu, Chiu-Chun
 10 TITLE OF INVENTION: HUMAN N-METHYL-D-ASPARTATE RECEPTOR
 11 TITLE OF INVENTION: SUBUNITS, NUCLEIC ACIDS ENCODING SAME AND USES THEREOF
 12 NUMBER OF SEQUENCES: 64
 13 ADDRESS/INVENTOR ADDRESS:
 14 ADDRESSEE: Heller Edman White & McCallan
 15 STREET: 4250 Executive Square, 7th Floor
 16 CITY: La Jolla
 17 STATE: CA
 18 COUNTRY: USA
 19 ZIP: 92037
 20 COMPUTER READABLE FORM:
 21 MEDIUM TYPE: Floppy disk
 22 COMPUTER: IBM PC compatible
 23 OPERATING SYSTEM: PC-DOS/MS-DOS
 24 SOFTWARE: Patent In Release #1.0, Version #1.25
 25 CURRENT APPLICATION DATA:
 26 APPLICATION NUMBER: US/08/940,086A
 27 FILING DATE: 29-SEP-97
 28 CLASSIFICATION: 546
 29 PRIOR APPLICATION DATA:
 30 APPLICATION NUMBER: US 08/231,193
 31 FILING DATE: 20-APR-1994
 32 CLASSIFICATION: 435
 33 PRIOR APPLICATION DATA:
 34 APPLICATION NUMBER: US 08/052,449
 35 FILING DATE: 20-APR-1994
 36 ATTORNEY/AGENT INFORMATION:
 37 NAME: Solomon, Stephanie
 38 REGISTRATION NUMBER: 44,779
 39 REFERENCE/PATENT NUMBER: 24745, 94860

GenCore version 4.5
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CM nucleic - nucleic search, using sw model

Run on: July 30, 2002, 02:07:10 ; Search time 1723.16 Seconds

(without alignments)
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File: US-09-932-678-1

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Total number of hits satisfying chosen parameters: 27472414

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Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	815.4	39.4	1800	10	Y16973 RNY16973 ka
2	675.4	32.7	788	10	HC502871 602550430
3	666	32.2	733	10	H1561523 603256222
4	653.6	31.6	996	10	BF203684 601866311
5	646	31.2	757	10	BE734354 601565566
6	637.4	30.8	1037	10	BE428305 602498983
7	617.8	29.9	671	9	AV703279 AV703279
8	614.6	29.7	736	10	H1464721 603202341
9	582.2	28.2	990	10	HG167061 602344740
10	553.4	26.6	828	10	H1761160 603043657
11	550.4	26.6	742	10	BEH70692 601448528
12	548	26.5	550	9	AL597171 DKEF26318K
13	547.2	26.5	727	10	B1549394 603190049
14	542.4	26.2	563	9	AW408066 U1-HP-BM0
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16	520.4	25.2	697	10	HG611364 602612725
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19	488.4	23.6	498	10	BE2110176
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24	470	22.7	490	10	BE502966
25	465.8	22.5	946	10	BE535886
26	453.2	22.4	632	10	BE614096
27	451.4	22.3	1045	10	H0247178
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29	451.4	21.8	461	9	A1928274
30	447.6	21.6	686	9	BE639379
31	440.4	21.3	956	10	BE295610
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35	421.8	20.4	547	9	AA530643
36	420.4	20.3	422	9	BE094077
37	419.2	20.3	811	10	BE560866
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39	406.4	19.7	479	9	AA239267
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ACCESSION	Y16973	1800 bp	mRNA	EST	Apr 1999
VERSION	Y16973	1800 bp	mRNA	EST	Apr 1999
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ORGANISM	Y16973	1800 bp	mRNA	EST	Apr 1999
REFERENCE	Y16973	1800 bp	mRNA	EST	Apr 1999
AUTHORS	Y16973	1800 bp	mRNA	EST	Apr 1999
TITLE	Y16973	1800 bp	mRNA	EST	Apr 1999
JOURNAL	Y16973	1800 bp	mRNA	EST	Apr 1999
COMMENT	Y16973	1800 bp	mRNA	EST	Apr 1999

FEATURES	Y16973	1800 bp	mRNA	EST	Apr 1999
Source	Y16973	1800 bp	mRNA	EST	Apr 1999
Location	Y16973	1800 bp	mRNA	EST	Apr 1999
Organism	Y16973	1800 bp	mRNA	EST	Apr 1999
Strain	Y16973	1800 bp	mRNA	EST	Apr 1999
Clone	Y16973	1800 bp	mRNA	EST	Apr 1999
Library	Y16973	1800 bp	mRNA	EST	Apr 1999
Vector	Y16973	1800 bp	mRNA	EST	Apr 1999
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RESULT	8
B1464721	
LOCUS	B1464721
DEFINITION	601202341P1 NTU_MGC_97 Homo sapiens cDNA clone IMAGE:5268011 5', mRNA sequence.

ACCESSION VERSION	KEYWORDS	SOURCE	ORGANISM	REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
B1464721	B1464721.1	G1:15255364 EST.	human.	Human sapiens	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.	1 (bases 1 to 736)	NIH-MGC http://mgc.nci.nih.gov/ .	National Institutes of Health, Mammalian Gene Collection (MGC) Unpublished (1999) Contact: Robert Strausberg, Ph.D.

TCGNA Library Preparation: Michael J. Brownstein (NHGRI), Shiroki Toshiyuki and Piero Carninci (RIKEN)
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MCC clone distribution information can be found through the I.M.A.G.E. Consortium/ILM at: <http://image.llnl.gov>
 Plate: ILAMI1676 row: b column: 12
 High quality sequence stop: 725.

FEATURES	LOCATION/Qualifiers
SOURCE	1. .736

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/clone_id="NH_MGC_97"
/lab_host="DH10B"
/note="Organ: testis; Vector: pBluescriptR (modified
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); Oligo-dt primed using primer 5'-TTTGTGTTTTTTTTVN-3'',
size-selected for average insert size 2.2 kb and
normalized to ROI 5. This is a primary library enriched
for full-length clones and constructed using the
Cap-trapper method (Carninci, in preparation). Library
constructed by M. Brownstein (NIMH/NICRI, National
Institutes of Health). Note: this is a NH_MGC library."

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ACCESSION	BC167061
VERSION	BC167061.1 GI:12674764
KEYWORDS	EST.
SOURCE	human.
ORGANISM	Homo sapiens
REFERENCE	Enkharjola; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo. 1 (bases 1 to 990)
AUTHOR	NIH-Mc7 http://mc7.nhl.nih.gov/ .
TITLE	National Institutes of Health, Mammalian Gene Collection (Mc7)

JOURNAL
COMMENT

Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: rstra@ncbi.nih.gov
Tissue Procurement: AFCE
cDNA Library Preparation: Ligo Technologies, Inc.
cDNA Library Arrayed by: The L.M.A.G.E. Consortium (LLNI)
RNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the L.M.A.G.E. Consortium/LLNI at:
http://image.llnl.gov

plate: LLAM0247 row: 9 column: 10
High quality sequence stop: 665.

FEATURES

Source

1. 990
/organism "Homo sapiens"
/db_xref "taxon:9606"
/clone "IMAGE:518495"
/clone_1lb "NHE_MGC_116"
/tissue_type "hypopharynx, cell line"
/lab_host "pH108 (phage resistant)"
/note "organ: kidney; Vector: pCMV-SPOK6; Site 1: Not:
Site 2: Salt: cloned unidirectionally; oligo-dT primed.
Average insert size 1.4 Kb. Library enriched for
full length clones and constructed by Ligo Technologies.
Note: This is a NIH-MGC Library."

BASE COUNT 604 a 275 c 219 g 192 t

ORIGIN

Query Match 28.8% Score 582.2; DB 10; Length 990;
Best Local Similarity 98.9% Prod. No. 6,46-155;
Matches 598; Conservative 0; Mismatches 8; Indels 1; Gaps 1;

1463 cgccttgggtttgacacagagcctttgacgagacatgaagagatggagta 1522

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1465 cgccttgggtttgacacagagcctttgacgagacatgaagagatggagta 1522

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551 llecysfhrAsnProLeuAspThrPhePheProPheAspProCysValLe 567
 1690 ATCTGACAAACCCGCTGGACACCTCTCTCCCTTGATGCCGCTGTGCT 1739
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seq_documentation_block:

ID AACT6937 standard: cDNA: 1418 BP.

AC AACT6937:

DT 08-FEB-2001 (first entry)

DE Human ORFX ORF2492 polynucleotide sequence SEQ ID NO:4981.

Human: open reading frame; ORFX; detection; cytosolic; hepatotropic;
 antiviral; antiparasitic; antiparkinsonian; neurotrophic; neuroprotective;
 anticonvulsant; osteoporotic; antiarthritic; immunosuppressant; cardiac;
 immunostimulant; thrombolytic; coagulant; vasotropic; antidiabetic;
 hypotensive; dermatological; immunosuppressive; antiinflammatory;
 antiviral; antibacterial; antifungal; antineumatic; antituberc;
 antineoplastic; gene therapy; cancer; proliferative disorder; hypertension;
 neurodegenerative disorder; osteoarthritis; graft vs host disease;
 cardiovascular disease; diabetes mellitus; hypothyroidism; SCID; AIDS;
 cholesterol ester storage; systemic lupus erythematosus; infection;
 severe combined immunodeficiency; malaria; autoimmune disorder; asthma;
 allergy; aplastic anaemia; nocturnal haemoglobinuria; burn; wound;
 bone damage; cartilage damage; antinflammatory disease; coagulation;
 thrombosis; contraceptive; ss.

OS Homo sapiens.

PN MO200058473-A2.

XX 05-OCT-2000

PF 31-MAR-2000; 2000MO-US08621.

PR 31-MAR-1999; 9905-0127607.

PR 02-APR-1999; 9908-0127636.

PR 05-APR-1999; 9905-0127728.

PR 30-MAR-2000; 2000US-0540763.

PA (CURA-) CURAGEN CORP.

PI Shinketsu RA, Leach M;

DR WPI: 2000-602362/57.

DR P-PSDB: AAB42728.

XX Novel nucleic acids and peptides derived from open reading frame X.
 PT useful for treating e.g. cancers, proliferative disorders,
 PT neurodegenerative disorders and cardiovascular disease

PS Claim 5: Page 4157-4158; 5507pp; English.

CC AAC74446 to AAC77606 encode the proteins given in AAB44497,
 CC which represent the human ORFX open reading frames 1 to 3141. The ORFX
 CC sequences have activities such as: cytosolic; hepatotropic; antiviral;
 CC antiparasitic; antiparkinsonian; neurotrophic; neuroprotective;
 CC osteoporotic; anticonvulsant; antiarthritic; immunosuppressant;
 CC immunostimulant; cardiac; thrombolytic; coagulant; vasotropic;
 CC antidiabetic; hypotensive; dermatological; immunosuppressive;
 CC antiviral; antibacterial; antifungal; antineumatic; antituberc;
 CC antituberc; and antineumatic. The sequences can be used for determining
 CC the presence of or predisposition to, or preventing or treating
 CC pathological conditions associated with an ORFX-associated disorder. The
 CC nucleic acids can be used to express ORFX proteins in gene therapy.
 CC vectors. The proteins and nucleic acids may be used to treat cancers,
 CC proliferative disorders, neurodegenerative disorders, osteoarthritis,
 CC graft vs host disease, cardiovascular disease, diabetes mellitus,
 CC hypertension, hypothyroidism, cholesterol ester storage, systemic lupus
 CC erythematosus, severe combined immunodeficiency (SCID), AIDS, viral,
 CC bacterial or fungal infection, malaria, autoimmune disorders, asthma,
 CC allergies, aplastic anaemia, burns, wounds, bone and cartilage damage,
 CC nocturnal haemoglobinuria, antinflammatory disease; to enhance
 CC coagulation; to inhibit thrombosis; and as a contraceptive.

XX Seq: 1418 BP; 422 A; 275 C; 318 G; 403 T; 0 other;

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Quality: 2162.00 Length: 422
 Ratio: 5.185 Gaps: 1
 Percent Similarity: 98.815 Percent Identity: 97.867

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 53 AAGAGCAACATGCAATGATGATGATGATGATGATGATGATGATGAT 102
 235 yRPheProThrLeuArgHisIleLeuGluLeuIleIleGluLysLe 251
 103 ATTTCACACCTTGGAGCATGAAATTCGACGTAATATATGAAAAAT 152
 251 uLeuLysLeuAspValAspAlaSerArgGlnGlyIleGluAspAlaGln 268
 153 ACTCAAGTGGATGATGATGATGATGATGATGATGATGATGATGATG 202
 268 LThrAlaThrIleThrCysGlyGlyThrAspSerThrGlnGlyLeuPhe 284
 202 AAACAGCAACATCAACTTTTGGTGGAGCAATCCAGGCAACATGATTT 252
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 253 AATATGGATGAAGATGAAGAACTGACATGAAACAAAGCTATGATG 302
 301 uArgLeuAspGlnMetValHisProValAlaGluArgLeuAspIleGln 318
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501 GATGATTAATCTGAAATCTGAAATCTGAAATCTGAAATCTGAAATCTGAAATCT 518
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903 TCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 952
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518 TATCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 544
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953 ATATCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 1002
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545 GATGATTAATCTGAAATCTGAAATCTGAAATCTGAAATCTGAAATCTGAAATCT 551
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1003 TATCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 1052
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551 cGlyThrAspProLeuAspThrIleThrIleThrIleThrIleThrIleThrIle 568
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1053 TCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 1102
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568 TATCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 584
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1103 TATCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 1152
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585 TCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 601
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1153 TATCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 1202
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601 TCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 618
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AC: AA501562:
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PE: 18-JUL-2001 (first entry)
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DE: Human secretory molecule. cDNA spm #52.
XX
KW: Human secretory molecule. cDNA spm #52.
XX
KW: cell signaling; cell proliferative disorder; attherosclerosis; cancer;
KW: immune system disorder; AIDS; neurological disorder; Alzheimer's disease;
KW: nervous system disorder; mental retardation; developmental disorder;
KW: neuromuscular disorder; microarray; Incyte ID number 428742ddec; SS.
XX
OS: Homo sapiens.
XX
PN: W0200123558-A2.
XX
PD: 05-Apr-2001.
XX
PE: 19-SEP-2000; 2000W0-0525610.
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PR: 28-SEP-1999; 990S-0156624.
PR: 28-SEP-1999; 990S-0156625.
PR: 02-DEC-1999; 990S-0168611.
PR: 02-DEC-1999; 990S-0168613.
PR: 02-DEC-1999; 990S-0168614.
XX
FA: (INV.) INYTE GENMUTS INV.
XX
PI: Hodgson DM, Lincoln SE, Russo FB, Spito PA, Banville SF,
PI: Bruchner SK, Duntour GE, Cohen HL, Rosen RH, Shah P, Chahap MS,
PI: Hillman JL, Jones AL, Yu JY, Greenwell LB, Panzer SK,
PI: Roseberry AM, Wladis RJ, Chen W, Liu TF, Yap PE, Stockbrecher JK,
PI: Anshov S. Pmid 11741.
OR: W01: 2001 258134/26.
XX
PT: New secretory polynucleotides (spm) and the polypeptides they encode,
PT: for use in inducing antibodies and screening libraries of compounds
XX
PS:
XX
CL: 1: Page 152; 161pp; English.
XX
CC: The present sequence for human secretory molecule cDNA spm #52
CC: (Incyte ID number 428742ddec) is 1 of 63 novel spm cDNA sequences
CC: (AA501511-AA501574) which encode for the secretory polypeptides spm. The
CC: spm polynucleotides are useful for screening a compound for
CC: effectiveness in altering expression of a target polynucleotide, where
CC: the target polynucleotide comprises spm. Spm is also useful in a method
CC: for assessing the toxicity of a test compound. Spm and its fragments or
CC: complementary sequences, may be used to identify the presence of and/or
CC: determine the degree of similarity between two nucleic acid sequences.
CC: Spm can also be used for a variety of diagnostic and therapeutic
CC: purposes, e.g., diagnosing a particular condition, disease or disorder
CC: associated with cell signaling, such conditions include cell
CC: proliferative disorders such as atherosclerosis, and cancers including
CC: leukemia, an immune system disorder e.g. acquired immunodeficiency
CC: syndrome (AIDS), a neurological disorder such as epilepsy or Alzheimer's
CC: disease, nutritional and metabolic disease of the nervous system, mental
CC: retardation and other developmental disorders, and muscular dystrophy and
CC: other neuromuscular disorders. Spm can also be used to design probes
CC: useful in diagnostic assays, which may be used to monitor the progress of
CC: conditions or disorders associated with abnormal levels of expression of
CC: spm. In addition spm encoding spm may be used for somatic or germline
CC: gene therapy, for inducing antibodies, or in microarrays.
XX
SQ: Sequence 1423 bp; 417 A; 247 C; 420 G; 439 T; 0 other:

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alignment_scores:

Quality: 1973.00 Length: 488
 Ratio: 4.995 Gaps: 2
 Percent Similarity: 80.943 Percent Identity: 80.533

alignment_block:

US-09-932-678-2 x AAS01562 ..

Align seq 1/1 to: AAS01562 from: 1 to: 1423

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47 q1ysThrValArgPheGlyGlyThrValThrGluValLeuLeuLysTyr 64
244 AAAAAGCTGTCGGTTGGTGGAAATGTGACAGAAAGTCTCTGAGTACA 293
64 y1ysGlyGlyIuThrAsnAspPheGluLeuLeuLysAsnGlnLeuLeuAsp 80
294 AAAAGCGTGAACAAATGACTTGGAGTTGTTGAAGAACCGCTGTTAGAT 343
81 ProAspIleIysAspAspGlnIleIleAsnTrpLeuLeuGluPheArgSe 97
344 CCACACATAAAGCATGACAGATCATCACTGGCTGCTAGCAATTCCTTC 394
97 rSerIleMetTyrLeuThrLysAspPheGluGlnLeuIleSerIleLeu 114
394 TTCTATATGTTACTTGCACAAAAGCTTGGACCACTTATCAGTATTATAT 443
114 euArgLeuProTrpIleuAsnArgSerGlnThrValValGluGluTyrLeu 130
444 TAAGATTGCTGGTGGTTCATAGCAAGTCAAAACAGTACAGCAAGATATTG 493
131 AlaPheLeuGlyLysLeuValSerAlaGlnThrValPheLeuAspProGly 147
494 GCTTTCTGCTGTAATCTTGATACGACAGACTGTTTCCACAGCCGTC 543
147 r1euSerMetIleAlaSerHisPheValProProArgValIleIleLysG 164
544 TCTGAGCATATGCTTCCCATTTTGCTCTCCCGAGTATGATTTAGG 593
164 IuGlyAspValAspValSerAspSerAspAspGluAspAspAsnLeuPro 180
594 AAGCGCATGTAGATGTTTCAGATTCTGATGATGATGATGATTAATCTTCT 643
181 AlaAsnPheAspThrCysHisArgAlaLeuGlnIleIleAlaArgTyrVal 197
644 GCAAAATTTGACACATGTCACAGACGCTTGCACAAATATAGCAGATATGT 693
197 rProSerThrProTrpPheLeuMetProIleLeuValGluLysPhePro 214
694 AGCATGCAACACGTTGTTCTCATGCAATACGTTGTCACAAAATTTTCAT 743
214 heValArgLysSerGluArgThrLeuGlnCysTyrValHisAsnLeuLeu 230
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447 heCysAspValAlaLeuHisGlyProPheTyrSerAlaCysGlnAlaVal 453
1174 TCTGCAATGTTGCTTTCATGAGCAATTTTACTGAGCTGCAAGCTGTCG 1223
444 PheTyrThrPheValPheArgHisLysGlnLeuLeuSerGlyLysLeuLeu 480
1224 TTCTACACCTTTGTTTTCACACAGCAAGCACTTTTCAAGCGAAACCTGAA 1273
480 sGluGlyLeuGlnTyrLeuGlnSerLeuAsnPheGluArgIleValMetS 497
1274 ACAAGGTTTGCAGATCTGAGATCTGAAATTTTGAGCGGATGATGATGA 1323
497 argGlnLeuAsnProLeuLysIleCysLeuProSerValValAsnProPhe 513
1324 GTCAGCTAAATGCCGTGAAGATTTGCTGCTGCAATGATTAATTTT 1373
514 AlaAlaIleThr 517
1374 GCTGCAATCACA 1385
seq_name: /s1db1/genedata/geneseq/geneseqn-emb:/NA2001A.DAT:AA544584
seq_documentation_block:
ID AA544584 standard; DNA: 1498 bp.
AC AA544584:
AT 18-DEC-2001 (first entry)
DT
XX
XX Human full-length polynucleotide sequence #9.
KW
KW Mammal: human; rhesus monkey; baker's yeast; fission yeast; Norway rat;
KW mouse; Chinese hamster; African clawed frog; fruit fly; dog; leek;
KW cancer; lymphoma; neuroblastoma; autoimmune disorder; cell proliferation;
KW nervous system disorder; inflammatory disorder; cell differentiation; ds;
KW aneuploidy; stem cell growth factor; actinin; inhibin; catalase; burn;
KW genetic disorder; bone regeneration; tendon; ligament; tissue repair;
KW

```



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67  uThrAspPheGluLeuLeuLysAsnGluLeuLeuAspProAspIleIle 84
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201 AACAAATGACTTGAAGTTGTTGAGAACAGCTGTAGATCCAGAACATAA 250
84  ys.....AspAspGluIleIleAsnProLeuGluIlePheArgSerSer 98
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251 AAGTGGTGTGCGCGCATGCCATATCTGTAGAGGAGGAGGAGGAGCAT 300
99  Ile.....MetTyrLeuThrLysAs 105
   ||| ||| |||
301 GTGGCCAGTGTGCTTTCATCCCTTGTGCTGTGCTGTGCTGTGCTGAAA 350
105 PheGluGluLeuIleSerIleIleLeuArgLeuProTyrPheLysArg 122
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351 CCAAGAGTGGCTCAGCCGAGAGGTGTGTAGCTGCTGCTGCTGCTGCTG 399
122 eTGIThrValIleGluIleTyrLeuIlePheLeuGluLysLeuValSer 138
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399 ..... 399
139 AlacIleThrValPheLeuArgProCysLeuSerMetIleAlaSerHis 155
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400 .....CCGCTGGCTGTGCGCCACATACAGCCCGCTGCTGCTGCTG 441
155 eValProTyrValIleIleLysGluGluLysValAspValSerAsp 172
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441 ..... 441
172 eTAspAspGluAspAspAsnLeuProAlaAsnPheAspThrCysHis 188
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442 .....AACCCCAACGCCACAGCTGTGCTGCTGCTGCTGCTGCTG 478
188 gAlaLeuGluIleIleAlaArgTyrValProSerThrProTyrPheLeu 205
   ||| ||| |||
479 .....CTGCTTGCAGG.....ACACCGCTGTCTCA 506
205 eTProIleLeuValGluLysPheProPheValArgLysSerGluArgThr 221
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507 TGGCAATAGCTGTGGAATAATTCATTTGTTCAAATACAGAGAACACA 556
222 LeuGluCysTyrValHisAsnLeuLeuArgIleSerValTyrPheProTh 238
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557 CTGCAATAGCTGTGCAATCTACTAGCATTTAGTGTATATTTTGCAC 606
238 rLeuArgHisGluIleLeuGluLeuIleIleGluLysLeuLeuLysLeu 255
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607 CTGAGCGCATGAATCTGCGAGCTTATTTGAAAACTACTCACTTGC 656
255 spValAsnAlaSerArgLysGlyIleGluLysPheAlaGluIleThrAlaThr 271
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657 ATGTGAATGATCCGGCAGGATTTGAGATGCTGAGAAACGCAAT 706
272 GluThrCysGlyIleThrAspSerThrGluGlyLeuPheAsnMetAspG 288
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707 CAACCTTGTGTGCGACAGATTCACGGAAGGATTCGTATATATGCTTAC 756
288 uAspGluGluThrGluHisGluThrLysAlaGlyProGluArgLeuAsp 305
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757 C..... 757
305 lMetValHisProValAlaGluArgLeuAspIleLeuMetSerLeuVal 321
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758 .....AGTTATTATGAAGTGGAG 778
322 LeuSerTyrMetLysAspValCysTyrValAspGlyLysValAspAsnG 338
   |||||
779 ATGAGTTTATC..... 790
338 yLysThrLysAspLeuTyrArgAspLeuIleAsnIlePheAspLysLeu 355
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791 .....ATAATCAAGAGTGGCAACACAGCTAG 815

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355 eTLeuProThrHisAlaSerCysHisValIlePhePheMetPheTyrLeu 371
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816 TGTGT..... 820
372 CysSerPheLysLeuGlyPheAlaValAlaPheLeuGluHisLeuPly 388
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821 .....CTATCTTTCATTCGACAGCCATTTTTCGAACTCTTTGAAA 864
388 sLysLeuGluAspProSerAsnProAlaIleIleArgValAlaValGly 404
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865 AAACCTGCAAGATTCACAGTAACTTCATCATCATCAGCGAGGTGTGGA 914
405 AsnTyrIleCysSerPheLeuAlaArgValLysPheIleProLeuLeu 421
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915 CATTAATATGCAAGCTTTTTCGACAGCTACATTAATTTCTCTTATAC 964
421 rValLysSerCysLeuAspLeuValAlaAsnTyrPheHisIleTyrLeu 438
   |||||
965 TGTAAACCATGGCTAGCTTTTGTGTAGCTGGCTGCACATATACCTTA 1014
438 sAsnGluAspSerGlyThrLysAlaPheCysAspValAlaLeuHisG 454
   |||||
1015 ATAACTCAGATTCGCGGAAACAAAGGCAATCTCGATGTGCTTCCATG 1064
454 lVProPheTyrSerAlaCysGlnAlaValPheTyrThrPheValPhe 470
   |||||
1065 GACCCATTTTACTGAGCTGTGCGCAAGCTGTGTCTGACCTTGTGTTTA 1114
470 rGHisLysGluLeuLeuSerGlyAsnLeuLysGluGlyLeuGluTyr 485
   |||||
1115 GAACACACAGACGCTTTTGAACGGAACCTGAAGAAGAGGCTGTGTTT 1172
seq_name: /STDB/ucvdat4/geneseq/geneseq-emb1/M2001B.DAI:AMS7646
seq_description_block:
ID AMS7646 standard: cDNA, 1017 bp.
XX
AC AA87636;
XX
DT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #23440.
XX
KW Human; Chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WC200175067-A2.
XX
PD 11-OCT-2001.
XX
PE 30 MAR-2001; 2001WO-050631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 24-AUG-2000; 2000US-0649167.
XX
PA (HENSE-) HYSEO INT.
XX
PI Diagnostic kit, Liu C, Tang YL;
XX
DK WPI: 2001-639162/73.
DK P-PSDB: ABG23449.
XX
New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity.
XX
PS Claim 1: SEQ ID No 23440: 104pp: English.
XX
CC The invention relates to isolated polynucleotide (1) and
CC polypeptide (11) sequences. (1) is useful as hybridisation probes.

```


seq documentation block:

ID ABA83038 standard: DNA: 1461 BP.

AC ABA83038;

DT 05-FEB-2002 (first entry)

DE Human transcription factor TRFX-65 coding sequence.

KW Human; transcription factor; TRFX; cell proliferative disease;

KW autoimmune disease; inflammation; neurological disease;

KW developmental disorder; cancer; AIDS; infection; cytosolic; anti-HIV;

KW neuroprotective; antiinflammatory; gene therapy; ds.

OS Homo sapiens.

PN W020017277-A2.

PD 04-OCT-2001.

PE 13-MAR-2001: 2001MO-US08117.

PR 13-MAR-2000: 2000OS-0188986.

PA (INCY-) INCYTE GENOMICS INC.

PI Hillman JL, Baughn MK, Yue H, Lai P, Lu DM, Patterson C;

PI Azimzai Y, Bandman O, Tang YT, Mathur P, Shah P, Au-Young J;

PI Reddy K;

DR WPI: 2001-570896/64.

DR P-PSDB: ABB50214.

PT Novel transcription factor polypeptides, used to treat diseases

PT associated with altered activity and expression of TRFX, and to screen

PT for agents capable of modulating its activity -

PS Claim 11: Page 299; 327pp: English.

XX The present sequence is the coding sequence for a human transcription

XX factor. The transcription factor and its coding sequence are useful in

XX the diagnosis, treatment and prevention of diseases associated with

XX altered expression of the transcription factor e.g. cell proliferative,

XX autoimmune/inflammatory, neurological and developmental disorders. A

XX number of specific disorders/diseases are given in the specification,

XX including: arteriosclerosis, cirrhosis, hepatitis, cancers, AIDS,

XX allergies, anaemia, asthma, autoimmune thyroiditis, bronchitis, atopic

XX dermatitis, diabetes mellitus, emphysema, Goodpasture's syndrome, gout,

XX Grave's disease, multiple sclerosis, osteoarthritis, pancreatitis,

XX psoriasis, rheumatoid arthritis, systemic lupus erythematosus, ulcerative

XX colitis, uveitis, Alzheimer's disease, Huntington's disease, Parkinson's

XX disease, stroke, and viral, bacterial, fungal and protozoal infections.

XX Sequence 1461 BP; 413 A; 445 C; 298 G; 415 T; 0 other:

alignment_scores:

Quality: 1124.00 Length: 486

Ratio: 3.986 Gaps: 8

Percent Similarity: 58.025 Percent Identity: 51.646

alignment_block:

US-09-932-678-2 x ABA83038

Align seq 1/1 to: ABA83038 from: 1 to: 1461

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195 ArgIyValIProSerThrProPheLeuMetProIleuValGluIy 211
    ::::|||||:|||||:|||||:|||||:|||||:|||||:|||||
263 AATATCAAAATTAAGGACCGCTGCTTCATGCCAATACTGCTGCAAA 312
    :|||||:|||||:|||||:|||||:|||||:|||||:|||||
211 spherophoeValIArGlySerGIuArFThLeuGIuYsYrValHSA 228
    |||||:|||||:|||||:|||||:|||||:|||||:|||||

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313 ATTCCATTGTCGAAATACAGAGAAACACTGCAATCTTACTTCATA 362
228 salLeuValArqIIeSerValITyPheProThIeUArqHISGIIuIleU 244
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363 ACTTACTAAGGATTACTGATATATTTCGCAACCTTCAGGCATGAAATTCG 412
245 GIuMetIleIerThIeULeuLysIleuValAsuAlaSerPArgI 261
    |||||:|||||:|||||:|||||:|||||:|||||:|||||
413 GAGTATTATTCGAAACACTACTGACCTGATGCTCAATGCAATCGGCA 462
261 nCIyIIeGIAsPAlaGluGluThIraIarGInIreCysGIyTha 278
    |||||:|||||:|||||:|||||:|||||:|||||:|||||
463 GGGATTTGAAGATGCTGTAACAAACATCAAAATCAAACTTGAGGGCAAC 512
278 sPserThrGIuIleuPheAsuMetAspCIuAspCIuGluHIS 294
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513 ATTCACGCAAGGATTCCTTTAATATG..... 538
295 GIuThIeUAlaGIuProGIuArGleuAspImetValIHISProVal 311
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311 aGIuArGIuAspIIeLeuMetSerLeuValIeUSeTYrMetLysAsp 328
    ..... 538
536 ..... 538
328 aICysTYrValAspGIyIleValAspAsuCIyIleThIeUAspIleYr 344
538 ..... 538
345 ArGAspIleuIleAsuIIePheAspIleuLeuIleuThIeUAlaSe 361
538 ..... 538
361 rCysHISValGIuPhePheMetPheTYrIleuCysSerPheIleuGIyP 378
539 .....GGAT 542
378 hAlaGIuAlaPheGluGluHISLeuTrpIleLysIleuGInAspProSe 394
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543 TCGACAGGCAATTTTCGACATCTTGGCAAAACCTTGAGATCCGACAT 592
395 AsPProAlaIleIleArGInAlaIaGIyAsuTYrIleCysSerPheP 411
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593 ATTCCTCCATCATACGCGAGCTCTCGAAATTAATTCGACACTTTT 642
411 uAlaArAlaIleSPheIIeProLeuIleTrValIleYSerCysIleuAsp 438
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643 GCGACAGGCTAAATTTATTTCTCTTATTTACTGTAAAGCATGCTATATC 692
438 eUleuValAsuTrpIleuHISIleTYrIleuAsuAsuIleAspSerGIyTh 444
    |||||:|||||:|||||:|||||:|||||:|||||:|||||
693 TTTGCTTAACTGGCTCGACATATACTTAATTAACGACATTCGCAACA 742
445 IysAlaPheCysAspValAlaIeHISGIyProPheTYrSerAlaYsGI 461
    |||||:|||||:|||||:|||||:|||||:|||||:|||||
745 AAGCATTCGCAATGCTCTCCATGAGCAATTTTACTTCAGCTCGCA 792
461 dAlaValIPheIyThrPheValIPheArqHISLysGIuIleuSeGIyA 478
    |||||:|||||:|||||:|||||:|||||:|||||:|||||
793 AATGTGTTTACAGCTTTGTGTTTATACACAAAGGAGCTTTTACAGGAA 842
478 salLeuYsGIuGIyIleuGInTYrIleuGInSerLeuAsuPheGIuArq 494
    |||||:|||||:|||||:|||||:|||||:|||||:|||||
843 ACGTGAAGCAAGCTTTCAGATATCTCAGATCTGAAATTTTGAAGGATA 892
495 ValMetSerGInIleuAsuProLeuLysIIeCysIleuProSerValValas 511
    |||||:|||||:|||||:|||||:|||||:|||||:|||||
893 GIGATGAGCCAGCCTAAATCCCTCGAAGATTTCCTGCTCGCTCAGTGTAA 942
511 nPhePheAlaAlaIleThrAsuLysTYrGIuIleuValIPheCysTYrThI 528
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943 CTTTTGTGCAATCACAAG..... 964

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560 PheProPhe AspProCysValLeuLysArgSerLysLysPhe IleAsp 575
 401 TTCCTCTTGATCCCTGTGTGCTGGAAGAGGTCAAGAAATTCATTCAT 450
 576 ProIle-Tyr-GlnValTyr-GluAspMetSerAlaIleGluLeuGlnI 591
 451 CCTATTTTATCCAGGTATGGGAGACATGATCTCTCAAGACGCTACAGCA 500
 591 uPheLysLysProMetLysLysAspIleValGluAspGluAspAsp 608
 501 GTTCAGAACCCATGAAAGAGCATAGTCAGACATGACATGATGATGACI 550
 608 heLeuLysGlyIleValProGlnAspAspThrValIleGlyIleThrPro 624
 551 TTTGGAAGGGAGAGTCCCGAGATGATACCGTATGGATGACACCA 600
 625 SerSerPheAspThrHisPheArgSerProSerSerSerValGlySerPr 641
 601 AGTCCTTTACAGCATTTCCGAGACTCTTCAAGTAGTGTGGCTGCCG 650
 641 oProValLeuTyr-MetGlnProSerProLeu 651
 651 ACCGCTCTGTACACGTGCAACCCAGTCCCTC 682
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seq_documentation_block:
 ID AA592253 standard; cDNA: 776 BP.
 AC AA592253;
 XX
 DT 13-FEB-2002 (first entry)
 XX
 DE DNA encoding novel human diagnostic protein #28057.
 XX
 KM Human: chromosome mapping; gene mapping; gene therapy; forensic;
 XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 OS Homo sapiens.
 XX
 PN W0200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PF 30-MAR-2001; 2001WO-0508631.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 PI Drmanac RT, Liu C, Tang YT;
 XX
 DR WPI: 2001-639462/73.
 DR P-PSDB; AB028066.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity
 XX
 PS claim 1: SEQ ID No 28057; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as

CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AA564197-AA594564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WPI,
 CC atftp.wpi.int/pub/published_pcl_sequences
 XX
 SO Sequence 776 BP; 206 A; 157 C; 198 G; 215 T; 0 other;

alignment_scores:
 Quality: 862.00 Length: 196
 Ratio: 4.717 Gaps: 5
 Percent Similarity: 95.408 Percent Identity: 94.898

alignment_block:
 US-09-932-678-2 x AA592253

Align seq 1/1 to: AA592253 from: 1 to: 776

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 192 ATGCGGCGACGCTCTGCTTACACGCTGTTGTGAGAGATCGAAGAC 247
 17 cSerSerAla-ValLysLysLeuGlyAlaSer ArgThrGlyIleSerAs 33
 248 CGCTCTTGCGTGAAGACGCTGTGCGGCGTGGAGCTGGGATTTGAAA 297
 33 uMetArgAlaLeuGluAsnAspPheAsnSerProThrArgLysThr 50
 298 TATCGGCGCATATACAGATCATTTTTCATTTCTGCGCCACAGAAAACTG 347
 348 TCGGTTGGTGGTGAAGCTGTGAGTGTGAAAGACGCTGTAAATGCAACAA 397
 50 AlaArgPheGlyGlyThrValThrGluValLeuLeuLysTyrLysGly 66
 392 GAAACAAATGACCTTGAAGTGTGAAAGACGCTGTAAATGCAACAA 447
 67 GluThrAsnAspPheGluLeuLeuLysAsnGlnLeuLeuAspProAsp 83
 448 AAGAGATGACCGACATATTAAGTGGCTGCTAGCAATTCCTCTCTGT 497
 83 cLysAspAsp-GluIleIleAsnTyr-LeuGluGluThrArgSerSer 99
 55 cMetTyrLeuThrLysAspPheGluLeuLeuLysSerIleIleLeuVal 116
 498 GATGATGCTTCCCATTTTGTCTCTCCGACGATGATTAAGAAAGG 697
 116 cProPro-TyrLeuAsnArgSerGlnThrValValGluGluTyrLeuAla 142
 548 TCGCTTGGTGGAAAGACGTCAACAGTAGTGAAGATTTGGCTT 597
 142 cLeuGluValSerLeuValSerAlaGlnThrValPheLeuArgProLysLeu 149
 598 TCTTGTAATCTTCTATCAGCAGACAGACTGTTTCTGTAAAGCTGTCTA 647
 149 cMetIleAlaSerHisPheValProProArgValIleIleLysGlyIle 165
 648 GATGATGCTTCCCATTTTGTCTCTCCGACGATGATTAAGAAAGG 697
 165 AspValAspValSerAspSerAspAspGluAspAspAsnLeuProAlaAs 182
 698 CATGATATGTTTCAGATTGTGATGATGAAGATGAATATCTTCTGAAA 747
 182 cPheAspThrCysHisArgAlaLeuGln 191
 748 TTTGACACATGTCACAGACGCTTTCGAA 775

seq_name: /SHS1/genbank4/geneseq/homoseq.cmbl/MA2001B.DAT:AA87643

seq document block:

10) AA87643 standard: cDNA: 1260 bp.

XX AA87643

XX 13 FEB 2002 (first entry)

XX cDNA encoding novel human diagnostic protein #24447.

XX Human chromosome mapping: gene mapping; gene therapy; forensic;

XX food supplement; medical imaging; diagnostics; genetic disorders; ss.

XX Homo sapiens.

XX W0200175067-A2.

XX 11 OCT 2001.

XX 60 MAR 2001: 2001WO-0508641.

XX 41 MAR 2000: 2000US-0540217.

XX 23 AUG 2000: 2000US-0649167.

XX (HENSE) HENSE INC.

XX Immune Kf, Liu C, Tang YF;

XX WPI: 2001-5-9362773.

XX P USDB: A062456.

XX New isolated polynucleotide and encoded polypeptides, useful in

XX diagnostics, forensics, gene mapping, identification of mutations

XX responsible for genetic disorders or other traits and to assess

XX biocompatibility

XX claim 1: SEQ ID No 24447: 104bp: English.

XX The invention relates to isolated polynucleotide (1) and

XX polypeptide (11) sequences, (1) is useful as hybridisation probes,

XX polymase chain reaction (PCR) primers, oligomers, and for chromosome

XX and gene mapping, and to recombinant production of (11). The

XX polynucleotides are also used in diagnostics as expressed sequence tags

XX for identifying expressed genes, (1) is useful in gene therapy techniques

XX to restore normal activity of (11) or to treat disease states involving

XX (11). (11) is useful for generating antibodies against it, detecting or

XX quantifying a polypeptide in tissue, as molecular weight markers and as

XX a food supplement. (11) and its binding partners are useful in medical

XX imaging of sites expressing (11). (1) and (11) are useful for treating

XX disorders involving abnormal protein expression or biological activity.

XX The polynucleotide and polynucleotide sequences have applications in

XX diagnostics, forensics, gene mapping, identification of mutations

XX responsible for genetic disorders or other traits to assess biodiversity

XX and to produce other types of data and products dependent on DNA and

XX amino acid sequences. AA87643/AA87644 represent novel human

XX diagnostic coding sequences of the invention.

XX Note: The sequence data for this patent did not appear in the printed

XX specification, but was obtained in electronic format directly from WPI

XX at http://wpi.int/pub/publihed-pat-sequences.

XX Sequence 1260 bp: 344 A: 227 C: 209 G: 420 T: 0 other:

XX alignment statistics:

XX Quality: 651.00 Length: 212

XX Pairs: 4.664 Gaps: 9

XX Percent Similarity: 91.981 Percent Identity: 91.038

XX alignment block:

XX US-09-932-678-2 X AA87643

Align seq 1/1 to: AA87643 from: 1 to: 1260

1 MetAlaAlaProLeuLeuHisThrArg-LeuProGlyAspAlaAlaAs 17

108 ATGGGAGGACAGGAGCTCTTCACACCTTTGTCGAGAGGACATGAGGCTT 247

17 GTCGTCATAGVALysylsLeuGlyAlaSer-MgProGlyIleSerAs 44

248 GTCGTCATAGVALysylsLeuGlyAlaSer-MgProGlyIleSerAs 297

33 GAGATGATAlaLeuGlyAsnAspHisPheAsnSerProProGlyIleV 50

298 TATGTCGATATAGAGAAATGATTTTAAATCTGCGCAAAAGAAAGAG 447

448 TTTGCTTTGAGTGGAGCTGTCGACAGCTTTCCTGACAGCTACAAAGAG 497

67 GTCATGAsnAspProGlyLeuLeuGlyAsnSerHisLeuLeuAspProAsp 84

498 GAAAGCAATGCACTTCAGCTTCAGAGAGAGAGAGAGAGAGAGAGAG 447

81 GlysAspSer-GlnIleLeuAsnTrp-LeuLeuGlyIleSerPro 99

448 AAGAGATGAGCTGATATATATATATATATATATATATATATATAT 497

99 GAGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 115

115 GAGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 141

548 ATTCGCTTTGCTGATGATGATGATGATGATGATGATGATGATGATG 597

141 GAGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 148

578 TTTCTTGTATGATGATGATGATGATGATGATGATGATGATGATG 647

148 GAGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 164

648 TTAGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 697

164 GAGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 181

698 GAGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 747

181 GAGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 197

748 GAGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 797

198 GAGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 204

798 GAGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 815

seq_name: /SHS1/genbank4/geneseq/homoseq.cmbl/MA2001B.DAT:AA87643

seq document block:

10) AA87643 standard: cDNA: 655 bp.

XX AA87643

XX 13 FEB 2002 (first entry)

XX cDNA encoding novel human diagnostic protein #16756.

XX Human chromosome mapping: gene mapping; gene therapy; forensic;

XX food supplement; medical imaging; diagnostics; genetic disorders; ss.

XX Homo sapiens.

XX W0200175067-A2.

XX 11 OCT 2001.

The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromosome and gene mapping, and in recombinant production of (II). The polynucleotides are also used in diagnostics as expressed sequence tags for identifying expressed genes. (I) is useful in gene therapy techniques to restore normal activity of (II) or to treat disease states involving (II). (II) is useful for generating antibodies against it, detecting or quantitating a polypeptide in tissue, as molecular weight markers and as a food supplement. (II) and its binding partners are useful in medical imaging of sites expressing (II). (I) and (II) are useful for treating disorders involving aberrant protein expression or biological activity. The polypeptide and polynucleotide sequences have applications in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits to assess biodiversity and to produce other types of data and products dependent on DNA and amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic coding sequences of the invention. Note: The sequence data for this patent did not appear in the printed specification, but was obtained in electronic format directly from WIPO at http://wipo.int/pub/published_pat_sequences.

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455 ATCCGACATAAAGCATGCACACTTCATCAATGGCTGTAGAAATTGG 406
96 qSerSerIleMetTyrLeuThrLysAspPheGlnGlnLeuIleSer11 112
405 TTCTTGCTGTAIGTACTGACCMAAACATTGAGCCACACTTATCAGTAT 356
112 cIlleGlnThrLeuProTPheGluAspArgScrGlnThrValValGluHis128
355 TATATTGAGATTCGCTTGTTGTGAATAGAAGTCAAAACAGTATGGAAG 306
129 TyrLeuAlaPheLeuGlyAsnLeuValSerAlaGlnThrValPheLeuAar 145
365 TATTGGCGCTTTCTTGGAATCTCTGTATCAGCATAGAGCTGTCTCTAG 256
145 qProcylauSerMetIleAlaSerHisPheValProProArgValIle 161
255 AACGATGTCACCATGATTTGCTTCTGCAATTTGTGCTCCXGXGAGTAG 206
162 IleySLGIUGLYASPVALASPIISERASPSORASPAAPGLIASPASAS 178
205 AITPAAGGAAGCGATGTABATTTTCACATTCGTGTGATGAAAGATGAA 156
178 nleuProAlaaspPheasPTHCysHisArgalaLeuGlnIleLeuVala 195
155 ICTTCTGCAAAATTTTGACACATATCACAGAGCTTGCAAAATAATAGCA 106
195 tgytYValProserThrProTrpPhe 203
105 GATATGTACCATCATGAGTATCTT 80
seq_name : /SIHSL/bcndata/geneseq/geneseqn-emb1/MAY1999.DAT:AAx51663
seq_documentation_block:
ID      AAx51663 standard; cDNA: 437 BP.
XX
XX      AAx51663:
XX
XX      21-JUN-1999 (first entry)
DE
DE      Human secreted protein 5' EST SEQ ID NO:242.
XX
XX      Human secreted protein; EST: expressed sequence tag; diagnosis;
KW      forensic; gene therapy; chromosome mapping; signal peptide;
UP      upstream regulatory sequence; cytokine activity; cell proliferation;
DI      differentiation; hemopoiesis regulation; tissue growth regulation;
KW      reproductive hormone regulation; chemotaxis; chemokinesis; haemostatic;
XX      thrombolytic; anti-inflammatory; tumour inhibition; ds.
XX
OS      Homo sapiens.
XX
XX      W05906549-A2.
XX
XX      PD 11-FEB-1999.
XX
XX      PF 31-JUN-1998: 98WC0-1401241
XX
XX      PR 01-AUG-1997: 97US-0905279.
XX
XX      (CEST ) GENSET.
XX
XX      PA Ductect A, Dumas Milne Edwards J., Lacroix B.;
PI
XX      WP1: 1999-154779/13.
XX      DR FJSDR: AAY12885.
XX
XX      NW nucleic acids encoding human secreted proteins - obtained from
PT      cDNA libraries derived from testis, ovary, uterus and spleen tissue
XX
XX      PS Claim 1: Page 344-345; 522pp; English.
XX
XX      AAx51459 to AAx1661 represent 5' expressed sequence tags (ESTs) for
XX      human secreted proteins, and encode the proteins given in AAT126d to
XX      AAT12913, respectively. The proteins given represent the signal peptide

```


0M 01: 05-09-932-678-2 to: N:geneseq_012802: * out_format : pts

Date: Jul 30, 2002 7:44 AM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:

-Model: fimer; p2n: model; -DEV: xih
-C/cq: 21/USPTO; spool: US09332678/rnatc_29072002_151516_19107/app-query.fasta.1.717
-DB: N:geneseq_032802 -Qfmt: fasta; -Suffix: oli; -mg: -GAP: 4.500
-GAPEXT: 0.050 -MINMATCH: 0.100 -XGAP: 0.000 -LOOPEXT: 0.000
-GAP: 4.500 -QGAPEXT: 0.050 -XGAP: 60.000 -XGAPEXT: 60.000
-GAP: 6.000 -GAPEXT: 7.000 -XGAP: 60.000 -XGAPEXT: 60.000
-DELCP: 6.000 -DELCPEXT: 7.000 -START: 1 -MATRIX: 01100
-TRANS: human40.cdt -LIST: 45 -DOCCALIGN: 200 -TRK_SCORE: quality
-THR_MIN: 1 -ALIGN: 15 -MODE: LOCAL -OUTFMT: pts -NORM: ext
-HEAPSIZE: 500 -MILEN: 0 -MAXLEN: 2000000000
-USER: 0509932678 -CGEN: 1.0 -ICPU: 6 -ICPU=3 -LONLOG
-DEV_TIMEOUT: 120 -WARN_TIMEOUT: 30 -NO_XLPXY -WAT -THREHOLD=1

Search information block:

Query: 05-09-932-678-2
Query length: 651
Database: N:Geneseq_032802: *
Database sequences: 1736436
Database length: 858457221
Search time (Sec): 212.800000

WARN: XGAP and XGAPEXT must be equal. Assuming XGAP=XGAPEXT=60.000
WARN: XGAPEXT and XGAPEXT must be equal. Assuming XGAPEXT=XGAPEXT=60.000

Score list:

Sequence	Strd Orig	ZScore	EScore	Len	Documentation
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/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA44584 +		295.00	5435.62	2.0e-294	1
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2000.DAT:AACT6947 +		276.00	5083.83	8.0e-275	14
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA501562 +		240.00	4416.46	1.2e-237	1
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA1999.DAT:AAV90248 +		97.00	1776.37	1.3e-90	34
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA87641 +		97.00	1759.62	1.2e-89	3
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA873660 +		93.00	1685.90	1.5e-85	2
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA891992 +		93.00	1685.90	1.5e-85	2
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA892253 +		79.00	1436.52	1.1e-71	7
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA1999.DAT:AA81565 +		75.00	1366.69	8.9e-68	4
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA892248 +		75.00	1364.44	1.2e-67	3
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA892254 +		71.00	1277.64	8.1e-63	3
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA883038 +		70.00	1264.93	4.1e-62	1
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA887636 +		69.00	1249.12	3.1e-61	1
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA882255 +		66.00	1187.02	9.0e-58	2
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA880954 +		63.00	1144.98	2.0e-55	3
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA887635 +		63.00	1144.98	2.0e-55	3
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA891996 +		63.00	1144.98	2.0e-55	3
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/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA885168 +		62.00	1092.92	1.6e-52	3
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2000.DAT:AACT7798 +		61.00	1095.36	1.2e-52	21
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2000.DAT:AA880955 +		59.00	1067.05	4.4e-51	4
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA887643 +		59.00	1062.13	8.2e-51	1
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/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA886764 +		58.00	1021.51	1.5e-48	2
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA885169 +		58.00	1021.51	1.5e-48	2
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:AA887640 +		54.00	974.65	6.1e-46	4
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/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2000.DAT:AACT7533 +		43.00	775.18	7.9e-35	34
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2000.DAT:AACT19091 +		38.00	686.21	7.1e-26	21
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA846215 +		35.00	625.60	1.7e-26	4
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA856760 +		35.00	625.60	1.7e-26	4
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA826382 +		35.00	625.60	1.7e-26	4
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AA84888 +		35.00	625.60	1.7e-26	4
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AAK30411 +		35.00	625.60	1.7e-26	4
/SIDSL/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:AAK115019 +		35.00	625.60	1.7e-26	4

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seq_documentation_block:
ID: AAAG8384 standard; cDNA: 2040 BP.
XX
XX
AC: AAAG8384;
XX
XX
UT: 08-FEB-2001 (first entry)
XX
XX
DE: Human RNA polymerase I transcription factor TIF-1A cDNA.
XX
XX
KW: RNA polymerase I transcription factor TIF-1A; antitumor; treatment;
KW: antiproliferative; cell proliferation; cancer; tissue regeneration; NS.
XX
XX
OS: Homo sapiens.
XX
XX
PN: W020055316-A1.
XX
XX
PO: 21-SEP-2000.
XX
XX
PE: 08-MAR-2000; 2000WD-DE00767.
XX
XX
PR: 17-MAR-1999; 99DE-1011992.
XX
XX
PA: (LRRK-) DEUT KRERSRKRSHONNSZHTNDOM.
XX
XX
PI: Giromit I, Winkler M;
XX
XX
WP: 2000-587527/55.
DR: P:SDS; AAB10946.
XX
XX
PT: New DNA encoding the transcription factor TIF-1A, useful for prevention
or treating diseases associated with abnormal cell proliferation,
particularly tumors.
XX
XX
PS: Claim 1; Fig 2; 38pp; German.
XX
XX
CC: This invention describes a novel DNA sequence (I) that encodes the RNA
polymerase I transcription factor TIF-1A which has antitumor,
CC: antiproliferative and proliferation-inducing activity. The invention also
describes (1) DNA (Ia) encoding a protein (II) with the biological
CC: activity of TIF-1A; (2) a ribozyme (R) corresponding to (I) or (Ia) and
able to bind specifically to, and cleave, its transcribed RNA so as to
CC: reduce or inhibit synthesis of the corresponding protein; (3) an
antisense RNA (AS) with binding properties similar to R; (4) an
CC: expression vector that contains (1), (Ia) or sequences that encode R or
AS; (5) host cells containing the vectors of (4); (6) TIF-1A or (II)
CC: encoded by (1) or (Ia); (7) preparation of TIF-1A or (II) by culturing
cells of (6); (8) ligands that bind to TIF-1A or (II); (9) antagonists
CC: that weaken or block the activity of TIF-1A or (II); (10) a diagnostic
method for detecting abnormal TIF-1A expression; and (11) kit for
CC: carrying out the method in (10). (1), (Ia) and similar sequences that encode
proteins with equivalent activity, expression vectors containing them, or
CC: their expression products are used to treat or prevent disorders
associated with reduced cellular proliferation, to stimulate cellular
proliferation, and to promote tissue regeneration, e.g. after injury or
CC: radiation therapy. Ribozymes, antisense sequences directed against (1),
also ligands and antagonists of TIF-1A, are used to treat or prevent (1),
CC: disorders associated with excessive cellular proliferation and to inhibit
proliferation, especially in treatment of cancers, (1) and specific
CC: ligands for TIF-1A (particularly antibodies (Ab)) are also useful for
diagnosis of altered TIF-1A expression by (indirect determination of the
CC: concentration, length and/or sequence of TIF-1A or its mRNA, e.g. for
detecting mutations). Ab can also be used for immunoprecipitation of
CC: TIF-1A and for isolation of related sequences from cDNA expression

[illegible]

XX	05-06T-20000.				
XX					
XX	41-MAR-2000: 2000060-0500621.				
XX					
XX	41-MAR-1999: 9908-0127607.				
XX	02-APR-1999: 9908-0127646.				
XX	05-APR-1999: 9908-0127728.				
XX	40-MAR-2000: 200005-0540764.				
XX					
XX	(70RA-) CUBACEN (COPR).				
XX					
XX	Shinkof's RA, Leach M;				
XX	WT: 2000 602462/57.				
XX	P-1506: AAB42728.				
XX					
XX	Move1 nucleic acids and peptides derived from open reading frame X,				
XX	useful for treating e.g. cancers, proliferative disorders,				
XX	neurodegenerative disorders and cardiovascular disease				
XX					
XX	Claim 5; Page 4157-4158; 5507pp; English.				
XX					
XX	AAC74446, to AAC77606 encode the proteins given in AAB40247 to AAB44497,				
XX	which represent the human ORFX open reading frames 1 to 4161. The ORFX				
XX	sequences have activities such as: cytosolic; hepatotropic; vitreoret;				
XX	antiproliferic; antiparkinsonian; modulatory; neuroprotective;				
XX	osteoplastic; anticoagulant; antithalitic; immunosuppressant;				
XX	immunostimulant; cardiant; thrombolytic; coagulant; vasodilator;				
XX	antidiabetic; hypotensive; dermatological; immunosuppressive;				
XX	antifibrotic; antibacterial; antiviral; antitumor; antihemetic;				
XX	antihypertoid; and antineurotic. The sequences can be used for determining				
XX	the presence of or predisposition to, or preventing or treating				
XX	pathological conditions associated with an ORFX-associated disorder. The				
XX	nucleic acids can be used to express ORFX proteins in gene therapy,				
XX	vectors. The proteins and nucleic acids may be used to treat cancers,				
XX	proliferative disorders, neurodegenerative disorders, osteoarthritis,				
XX	graft vs host disease, cardiovascular disease, diabetes mellitus,				
XX	hypercholesterolemia, hypohydrotism, cholesterol ester storage, systemic lupus				
XX	erythematosus, severe combined immunodeficiency (SCID), AIDS, viral,				
XX	hepatitis or fungal infection, malaria, autoimmune disorders, asthma,				
XX	allergies, aplastic anemia, lupus, wounds, bone and cartilage damage,				
XX	neutrophil hemophilinuria, antinflammatory disease, to enhance				
XX	coagulation; to inhibit thrombosis; and as a cell receptor.				
XX					
XX	Sequence 1418 MF: 422 A; 276 G; 418 G; 404 T; 0 other;				
XX					
XX	alignment_scores:				
XX	Quality: 276.00 Length: 276				
XX	Ratio: 1.000 Gaps: 0				
XX	Percent Similarity: 100.000 Percent Identity: 100.000				
XX					
XX	Alignment block:				
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XX					
XX	Align seq 1/1 for AAC76937 from: 1 to: 1418				
XX					
XX	276 GlycyllybAspSerThrGluGlySerbAsnMetAspGluAspGlu 291				
XX					
XX	225 GSHGRCAGATTCACGAGAGATTCCTTAAATATGATGAGAAATGAGAA 272				
XX	291 GSHGRCAGATTCACGAGAGATTCCTTAAATATGATGAGAAATGAGAA 308				
XX					
XX	274 AAATCAAAATCAAAATCAAAATCAAAATCAAAATCAAAATCAAAATCA 322				
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XX	308 TAPGVATATAGTATAGTATAGTATAGTATAGTATAGTATAGTATAGTAT 324				
XX					
XX	325 ATGCTATAGTATAGTATAGTATAGTATAGTATAGTATAGTATAGTAT 372				
XX					
XX	325 M-1TGSASVValTcysTValAspTcylgysValAspAsnGlyTcylly 341				
XX					
XX	47-AGGAGAGATTCGCTATGCTATGCTATGCTATGCTATGCTATGCTATG 422				


```

341 sasplpaurtrarasplleuileasullepheaspysleuleuileuprot 458
|||||
423 GATATATATCGGACCTGATATAACATCTTACACAACTCTCTGTCGCCA 472
358 hrhlsalaserCysHisValGlnphehemephetryleucysSerPhe 374
|||||
474 CCATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 522
375 LysLeuGlyPheAlaGlnAlaPheLeuGlnHisLeuTyrPheLysLeuG 391
524 AATTCGGATTCGCAAGGCACTTTGCAACAACCTGCGAATAATTCGA 572
391 naspprosoraspptolalleleparGlnAlaIleGlnAlaGlnAsnTyr 408
|||||
573 GACCCAACTATCTCTGCAATCAGGCGGCTGCTGCAATATATATG 622
408 LysSerPheLeuAlaArgAlaLysPheLeuProLeuIleThrValLysSer 424
|||||
623 GAACCTTTTGGCAAGACCAATATATCTCTGCTTATCTGCTAAATCA 672
425 CysLeuAspLeuLeuValAsnTripleuHisIleTyrLeuAsnAsnGlnAs 441
|||||
673 TCGCTAGACATCTTTGCTTAACCTGCTGCAATATCTGCTAAATCA 722
441 pserGlyThrLysAlaPheCysAspValAlaLeuHisGlyProPheTyrS 458
|||||
723 TTCGGCAACAAAGGCTTTGCAATGCTGCTGCTGCTGCTGCTGCTGCT 772
458 eValAcGlycylAlaValPheTyrThrPheValPheArgHisLysGlnLeu 474
|||||
774 CAGCCGCAAGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 822
475 LeuSerGlyAsnLeuLysGlnLysLeuGlnTyrLeuGlnSerLeuAsnPh 491
|||||
823 TGCAGGCAAGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 872
491 eGluArgIleValMetSerGlnLeuAsnProLeuLysIleGlySerProS 508
|||||
873 TACGCGGATGATGAGGAGGCTAAATCTGCTGCAATTTCTGCTGCT 922
508 eValValAsnPhPheAlaAlaIleThrAsnLysTyrGlnLeuValPhe 524
|||||
923 CAGTGTATACCTTTTCTGCTGCAATCACAATAAGTACAGCTGCTGCT 972
525 CysTyrThrIleLeuGlnArgAsnAsnArgGlnMetLeuProValIleAr 541
973 TGTACACCATCATGAGAGCAACATCGCAATGCTGCTGCTGCTGCTGCT 1022
541 gserThrAlaGlyGlyAspSerValGln 550
|||||
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seq_name: /SIUS1/gcdata/geneseq/geneseq-emb1/NA2001A.DAT:AA501562

seq_documentation_block:

ID: AA501562 standard; cDNA: 1423 BP.

AA501562:

18-JUL-2001 (first entry)

Human secretory molecule cDNA spm #52.

Human: secretory molecule; spm: SPTM; library screening; gene therapy;

cell signaling; cell proliferative disorder; atherosclerosis; cancer;

immune system disorder; AIDS; neurological disorder; Alzheimer's disease;

neurovascular disorder; mental retardation; developmental disorder;

neurovascular disorder; microarray; Inyte ID number 4287452dec; ss.

Humo sapiens.

XX OS

XX PN W0200123558-A2.

```

XX 05-APR-2001.
PD 19-SEP-2000: 2000W0-0525610.
XX 28-SEP-1999: 9905-0156624.
PR 28-SEP-1999: 9905-0156625.
PR 02-DEC-1999: 9905-0158611.
PR 02-DEC-1999: 9905-0158613.
PR 02-DEC-1999: 9905-0158614.
XX (INCY-) INCYTE GENOMICS, INC.
PA Hodgson DM, Lincoln SE, Russo FD, Spito PA, Banville SC,
PI Bratcher SR, Imour GE, Cohen BJ, Rosen BH, Shah P, Chalup MS,
PI Hillman JL, Jones AL, Yu JY, Greenwalt JB, Panzer SR,
PI Roseberry AM, Wright RJ, Chen W, Liu TF, Yap PE, Stockdreyer IK,
PI Anshey S, Fonj WJ.
XX WPI: 2001-258134/25.
DR
XX New secretory polynucleotides (spm) and the polypeptides they encode,
PT for use in inducing antibodies and screening libraries of compounds
XX
XX Claim 1: Page 152; 161pp; English.
PS
XX The present sequence for human secretory molecule cDNA spm #52
CC (Inyte ID number 4287452dec) is 1 of 63 novel spm cDNA sequences
CC (AA501511-AA501573) which encode for the secretory polypeptides spm. The
CC spm polynucleotides are useful for screening a compound for
CC effectiveness in altering expression of a target polynucleotide, where
CC the target polynucleotide comprises spm. spm is also useful in a method
CC for assessing the toxicity of a test compound. spm and its fragments or
CC complementary sequences, may be used to identify the presence of and/or
CC determine the degree of similarity between two nucleic acid sequences.
CC spm can also be used for a variety of diagnostic and therapeutic
CC purposes, e.g. diagnosis of a particular condition, disease or disorder
CC associated with cell signaling, such conditions include cell
CC proliferative disorders such as atherosclerosis, and cancers including
CC leukemia, an immune system disorder e.g. acquired immunodeficiency
CC syndrome (AIDS), a neurological disorder such as epilepsy or Alzheimer's
CC disease, nutritional and metabolic disorder of the nervous system, mental
CC retardation and other developmental disorders, and muscular dystrophy and
CC other neuromuscular disorders. spm can also be used to design probes
CC useful in diagnostic assays, which may be used to monitor the progress of
CC conditions or disorders associated with abnormal levels of expression of
CC spm. In addition spm encoding SPTM may be used for somatic or germ-line
CC gene therapy, for inducing antibodies, or in microarrays.
XX
XX Sequence 1423 bp: 417 A; 247 C; 320 G; 439 T; 0 other:

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alignment_scores:

Quality:	240.00	Length:	240
Ratio:	1.000	Gaps:	0
Percent Similarity:	100.000	Percent Identity:	100.000

alignment_block:

us-09-932-678-2 x AA501562

Align seq 1/1 to: AA501562 from: 1 to: 1423

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194 ATTTAAATATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 244
47 gserThrValArgPheGlyGlyThrValThrGlnValAlaLeuLysTyrL 64
|||||
244 AAAAAATGTTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 293
64 yslGlyGlyLthrAsnAspPheGlnLeuLeuLeuLysAsnGlnLeuAsp 80
|||||
293 AAAAAAGTGAACAAATATGACTTTGAGTTGTTGCAAGACAGCTGTTAAAT 343

```



```

XX  DNA encoding novel human diagnostic protein #23445.
DE
XX
XX  Human: chromosome mapping; gene mapping; gene therapy; forensic
KM
XX  food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
XX  Homo sapiens.
OS
XX  W0200175067-A2.
FN
XX  11-OCT-2001.
PD
XX
XX  30-MAR-2001; 2001WO-0508631.
FE
XX
XX  41-MAH-2000; 2000US-0540217.
PR
XX  23-AUG-2000; 2000US-0649167.
PR
XX
XX  (HYSE-) HYSEQ INC.
PA
XX
XX  Drmanac RT, Liu C, Tang YT.
P1
XX
XX  WPI: 2001-639362/73.
DK
XX  P-PSDB; AAC23445.
D8
XX
XX  New isolated polynucleotide and encoded polypeptides, useful in
PT  diagnostics, forensics, gene mapping, identification of mutations
PT  responsible for genetic disorders or other traits and to assess
PT  biodiversity -
XX
XX  (Claim 1; SEQ ID NO 23445; 103pp; English.
PS
XX
XX  The invention relates to isolated polynucleotide (I) and
CC  polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC  polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC  and gene mapping, and in recombinant production of (II). The
CC  polynucleotides are also used in diagnostics as expressed sequence tags
CC  for identifying expressed genes. (I) is useful in gene therapy techniques
CC  to restore normal activity of (II) or to treat disease states involving
CC  (II). (II) is useful for generating antibodies against it, detecting or
CC  quantitating a polypeptide in tissue, as molecular weight markers and as
CC  a food supplement. (II) and its binding partners are useful in medical
CC  imaging of sites expressing (II). (I) and (II) are useful for treating
CC  disorders involving aberrant protein expression or biochemical activity.
CC  The polypeptide and polynucleotide sequences have applications in
CC  diagnostics, forensics, gene mapping, identification of mutations
CC  responsible for genetic disorders or other traits to assess biodiversity
CC  and to produce other types of data and products dependent on DNA and
CC  amino acid sequences. AAC64197-AAS94564 represent novel human
CC  diagnostic coding sequences of the invention.
CC  Note: The sequence data for this patent did not appear in the printed
CC  specification, but was obtained in electronic format directly from Wipo
CC  at http://wipo.int/pub/published\_pat\_sequences.
XX
XX  Sequence 3169 BP; 812 A; 841 C; 779 G; 737 T; 0 other:
50

```

419 polr1b Valylsercysleupenleuvalasutprrwhstle 435
|||||
957 ttttttactgttataatcatgctgtgaaatttttttggttaacctggtaacata 1006
436 tttttaaanaaaactaaasppertlythetysalabheycasfvalala 452
|||||
1007 tacccttaataaacatgagcatctgggaaacaaagacatctggcatgttacct 1056
1057 ccattgacacaaatgaccttttgaaacggaaacctgaaagacacac 1147
469 hearrhsttststleuleusercylasnhleuylsttsttly 482
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1107 tttagacacaaatgaccttttgaaacggaaacctgaaagacacac 1147

seq_name: /SIDSL/ncdata/a/geneseq/geneseqn-emb1/NA2001b.fai:AA573660

seq_documentation block:
ID: AA573660 standard; cDNA; 2991 bp.
XX
AC: AA573660;
XX
DT: 13 FEB-2002 (first entry)
XX
DE: DNA encoding novel human diagnostic protein #9464.
XX
KW: Human; chromosome mapping; gene mapping; gene therapy; diagnostic;
KW: food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS: Homo sapiens.
XX
PN: W0200175067-A2.
XX
PD: 11 OCT-2001.
XX
PE: 30-MAR-2001; 2001W0-0508631.
XX
PR: 31 MAR-2000; 2000O05-0540217;
PR: 23 APR-2000; 2000O05-0649167.
XX
PA: (HSE-) HYSEQ INC.
XX
PI: Dmanac RT, Liu C, Tanq YI;
XX
DR: WP1: 2001-619362/773;
DR: P-PSDB; AEG09473.
XX
PT: New isolated polynucleotide and encoded polypeptides, useful in
PT: diagnostics, forensics, gene mapping, identification of mutations
PI: responsible for genetic disorders or other traits and to assess
PI: biodiversity -
XX
PS: Claim 1; SEQ ID NO 9464; 10pp; English.
XX
CC: The invention relates to isolated polynucleotide (I) and
CC: polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC: polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC: and gene mapping, and in recombinant production of (II). The
CC: polynucleotides are also used to diagnose as expressed sequence tags
CC: for identifying expressed genes. (I) is useful in gene therapy techniques
CC: to restore normal activity of (II) or to treat disease states involving
CC: (II). (II) is useful for generating antibodies against it, detecting or
CC: quantitating a polypeptide in tissue, as molecular weight markers and as
CC: a food supplement. (II) and its binding partners are useful in medical
CC: imaging of sites expressing (II). (I) and (II) are useful for treating
CC: disorders involving aberrant protein expression or biological activity.
CC: The polypeptide and polynucleotide sequences have applications in
CC: diagnostics, forensics, gene mapping, identification of mutations
CC: responsible for genetic disorders or other traits to assess biodiversity
CC: and to produce other types of data and products dependent on DNA and
CC: amino acid sequences. AA564197-NA594564 represent novel human


```

XX 13-FEB-2002 (first entry)
XX DNA encoding novel human diagnostic protein #28057.
DE
XX
XX Human: chromosome mapping; gene mapping; gene therapy; forensic:
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX Homo sapiens.
XX M0200175067-A2.
XX
XX 11-OCT-2001.
XX
XX 30-MAR-2001: 2001WO-US08631.
XX
XX 31-MAR-2000: 2000US-0540217.
XX 23-AUG-2000: 2000US-0649167.
XX
XX (HSE-) HSEQ INC.
XX
XX Drmanac RT, Liu C, Tang YT.
XX
XX WP1: 2001-63362/73.
XX P-PSDB; ABG28066.
XX
XX New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity.
XX
XX Claim 1: SEQ ID No 28057; 103pp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences, (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits to assess biodiversity
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. AAS64197-AAS94564 represent novel human
XX diagnostic coding sequences of the invention.
XX Note: The sequence data for this patent did not appear in the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pcl_sequences.
XX
XX Sequence 776 BP; 206 A; 157 C; 198 G; 215 T; 0 other:

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Alignment_scores:

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Quality: 79.00 Length: 93
Ratio: 0.859 Gaps: 1
Percent Similarity: 98.925 Percent Identity: 98.925

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alignment_block:

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US-09-932-678-2 x AAS92253 ..
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111
499 ATGACTTGACCAAGACCTTGAGCAACTATCATGATATATGAAATT 548
116 uPro.TripLeuAsnArgSerGlnThrValValGlnGluTyrIleuAlaIle 132

```

```

|||||
54- GCGTTGCTGCTGATAGACAGCTTAACACAGTAGTCGACAGCTATTGCGTTT 598
133 LeuGlyAsnLeuValSerAlaGlnThrValPheLeuAlaGlyProCysLeuSe 149
599 CTTCGAAATCTGCTGATGACAGACAGCTGCTTCTGACAGCGCTGCTGAG 648
149 TGGTTCGAAATCTGCTGATGACAGACAGCTGCTTCTGACAGCGCTGCTGAG 166
649 CAGGATTCGCTGCAATTTGCTGCTGCTGCAATTCATTAAGCAAGAGG 698
166 ValAlaSerValSerAspSerAspAspGluAspAspAsnLeuProAlaAsn 182
699 AGTACAGATCTTTCAGATCTTCATGATGAAAGATGATATCTTCCTCAAT 748
183 PheAspThrCysHisArgAlaLeuGln 191
749 TTGGACACATGCTGAGAGCGCTTGCAA 775
seq_name: /SI0SI/seqdata/geneseq/geneseq_emb1/NA1999.DAT:AA51663
seq_documentation_block:
ID AA51663 standard; cDNA; 437 BP.
XX
XX AA51663:
AC
XX
XX 21-JUN-1999 (first entry)
DE
XX Human secreted protein 5' EST SEQ ID NO:242.
XX
XX Howin: secreted protein; EST; expressed sequence tag; diagnostic;
XX forensic; gene therapy; chromosome mapping; signal peptide;
XX upstream regulatory sequence; cytokine activity; cell proliferation;
XX differentiation; hematopoiesis regulation; tissue growth regulation;
XX retroviral hormone regulation; chemokine; chemokine; haemostatic;
XX thrombolytic; anti-inflammatory; tumour inhibition; ds.
XX
XX Homo sapiens.
XX
XX M05905549-A2.
XX
XX 11-FEB-1999.
XX
XX 31-JUL-1998: 98WO-1801231.
XX
XX 01-AUG-1997: 97US-0905279.
XX
XX (CDS1 ) CDSSET.
XX
XX Dacfort A, Dumas Milne Edwards J, Lacroix B.
XX
XX WP1: 1999-154794/13.
XX P-PSDB; AAY12885.
XX
XX New nucleic acids encoding human secreted proteins obtained from
XX cDNA libraries derived from testis, ovary, uterus and spleen tissue
XX
XX Claim 1: Page 344-345; 522pp; English.
XX
XX AA51459 to AA51691 represent 5' expressed sequence tags (ESTs) for
XX human secreted proteins, and encode the proteins given in AAY12641 to
XX AAY12913, respectively. The proteins given represent the signal peptide
XX and an N-terminal fragment of a secreted protein. The nucleic acid
XX sequences can be used for producing secreted human gene products. They
XX can also be used to develop products for diagnosis and therapy. The
XX proteins obtained may have cytokine activity, cell
XX proliferation/differentiation activity, haematopoiesis regulation
XX activity, tissue growth regulating activity, reproductive hormone
XX regulating activity, chemokine/chemokine activity, haemostatic and
XX thrombolytic activity, receptor/ligand activity, anti-inflammatory
XX activity, tumour inhibition activity or other activities. The products
XX can be used in forensic, gene therapy and chromosome mapping procedures.
XX The sequences can also be used for obtaining corresponding promoter

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Hot! Saper's

XX WP0200175067.A2.
 XX
 XX 11-oct-2001.
 XX
 XX 10-MAR-2001: 2001MO-US08631.
 XX
 XX 31-MAR-2000: 2000US-0540217.
 XX 23-AUG-2000: 2000US-0649167.
 XX
 XX (HYSEQ-) HYSEQ INC.
 XX
 XX human CT, Liu C., Tang YT;
 XX
 XX WPI: 2001-639362/73.
 XX P-PSDB: ABG28067.
 XX
 XX Now isolated polynucleotide and encoded polypeptides, useful in
 XX diagnostics, forensics, gene mapping, identification of mutations
 XX responsible for genetic disorders or other traits and to assess
 XX biodiversity.
 XX
 XX Claim 1: SEQ ID No 28058; 103pp; English.
 XX
 XX The invention relates to isolated polynucleotide (I) and
 XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
 XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 XX CC and gene mapping, and in recombinant production of (II). The
 XX polynucleotides are also used in diagnostics as expressed sequence tags
 XX for identifying expressed genes. (I) is useful in gene therapy techniques
 XX CC to restore normal activity of (II) or to treat disease states involving
 XX (II). (II) is useful for generating antibodies against it, detecting or
 XX quantitating a polypeptide in tissue, as molecular weight markers and as
 XX a food supplement. (II) and its binding partners are useful in medical
 XX imaging of sites expressing (II). (I) and (II) are useful for treating
 XX disorders involving aberrant protein expression or biological activity.
 XX The polypeptide and polynucleotide sequences have applications in
 XX diagnostics, forensics, gene mapping, identification of mutations
 XX responsible for genetic disorders or other traits to assess biodiversity
 XX CC and to produce other types of data and products dependent on DNA and
 XX amino acid sequences. AAS64197-AAS94564 represent novel human
 XX diagnostic coding sequences of the invention.
 XX Note: The sequence data for this patent did not appear in the printed
 XX CC specification, but was obtained in electronic format directly from Wipo
 XX at http://wipo.int/pub/published_pat_sequences.
 XX
 XX Sequence 3175 BP: 811 A: 849 C: 779 G: 736 T: 0 other:
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 XX Percent Similarity: 100.000 Percent Identity: 100.000
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 XX US-09-932-678-2 x AAS92254 ..
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 XX Align seq 1/1 to: AAS92254 from: 1 to: 3175
 XX
 XX 200 ThrProTrrPheValuMetProIleuValGluLysPheProPheValAr 216
 XX ||||||||||||||||||||||||||||||||||||||||||||||||||
 XX 491 ACACCGTGGTTCATGCAATGCAATGCGAATAATTTCCATTGTCG 540
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 XX 216 gysserGluArGThrLeuGluGcSTyValHisAsnLeuLeuArGiles 233
 XX ||||||||||||||||||||||||||||||||||||||||||||||||||
 XX 541 AAAATACAGAGAGACACTGGAATGTACGCTTCATACCTTACTAAGCATTA 590
 XX ||||||||||||||||||||||||||||||||||||||||||||||||||
 XX 233 eValTyrrPheProThrLeuArGHisGluLLeuGluLeuLeuIleGlu 249
 XX ||||||||||||||||||||||||||||||||||||||||||||||||||
 XX 591 GGTGATATTTCCAACTTGAGCGATGAAATTCGGAGCTTATATTCGAA 640
 XX ||||||||||||||||||||||||||||||||||||||||||||||||||
 XX 250 LysIleuLeuLysLeuAspValAsnAlaSerArgGlnGlyIleGluAspAl 266
 XX ||||||||||||||||||||||||||||||||||||||||||||||||||

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266 actluc10tHuaAa_270
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641 TCAAGCAACACAA_703

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seq_documentation_block:
id ABA83038 standard: DNA; 1461 bp.
AC ABA83338:
XX
XX 05-FEB-2002 (first entry)
DT
DE Human transcription factor TRFX-65 coding sequence.
XX
XX Human transcription factor: TRFX: cell proliferative disease;
KW autoimmune disease; inflammation; neurological disease;
KM immunofund disorder; cancer; AIDS; infection; cytostatic anti HIV;
KW neuroprotective; antiinflammatory; gene therapy; ds.
XX
XX Homo sapiens.
OS
XX Wc200172777-A2.
XX
XX 04-OCT-2001.
PD
PE 13-MAR-2001: 2001W-US08117.
XX
XX 13-MAR-2000: 2000US-0189986.
PA
PA (INCYTE GENOMICS INC.
XX
XX Hillman JL, Raughn MR, Yue H, Lai P, Lu DM, Patterson CJ;
PI Azharai Y, Bandman O, Tang YT, Mathur P, Shah P, Au-Yang J;
PI Ready R;
XX
XX WP1: 2001-570896/64.
DR
DR P FSDS: ABB50214.
XX
XX Novel transcription factor polypeptides, used to treat diseases
PT associated with altered activity and expression of TRFX, and to screen
PT for agents capable of modulating its activity.
XX
XX Claim 11; Page 299; 327pp; English.
PS
XX

The present sequence is the coding sequence for a human transcription
factor. The transcription factor and its coding sequence are useful in
the diagnosis, treatment and prevention of diseases associated with
altered expression of the transcription factor e.g. cell proliferative,
autoimmune/inflammatory, neurological and developmental disorders. A
number of specific disorders/diseases are given in the specification,
including: arteriosclerosis, cirrhosis, hepatitis, cancers, AIDS,
allergies, anaemia, asthma, autoimmune thyroiditis, prothitis, atopic
dermatitis, diabetes mellitus, emphysema, Goodpasture's syndrome, gout,
Graves's disease, multiple sclerosis, osteoarthritis, psoriasis,
psoriasis, rheumatoid arthritis, systemic lupus erythematosus, ulcerative
colitis, uveitis, Alzheimer's disease, Huntington's disease, Parkinson's
disease, stroke, acid viral, bacterial, fungal and protozoal infections.
XX
XX Sequence 1461 BP; 413 A; 335 C; 298 G; 415 T; 0 other:

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Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
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Align seq 1/1 to: ABA83038 from: 1 to: 1461

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XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.

PS Claim 1: SEQ ID No 28059; 103bp; English.

XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at http://wipo.int/pub/published_prt_sequences.

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alignment_scores:
 Quality: 66.00 Length: 66
 Ratio: 1.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

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 1 GATCTTTGCTTAAGCTGCTGCACATATACCTTAATAACGAGATTGCGG 50
 443 yIhrlYsAlaPhcCysAspValAlaIleuHisGlyProPhcTYrSerAlaG 460
 ||||||||||||||||||||||||||||||||||||||||||||||||
 51 AACCAAGAGCATTCGTGATGTTGCTCTCATGACCATTTTACTCAGGCT 100
 460 ysgInAlaValPheTYrThrpheValPheArGHisLysGlnLeuLeuSer 476
 ||||||||||||||||||||||||||||||||||||||||||||||||
 101 GCCAACCTGTGTACACCTTTGTTTACAGACCAAGCAGCTTTGAAC 150
 477 G1yAsnLeuLysGlnGlyLeuGlnTYrLeuGlnSerLysAsnPhcGlu 492
 ||||||||||||||||||||||||||||||||||||||||||||||||
 151 GCAAACTGAAGAAGGTTTGCAGTAICTTCAAGACTGCAATTTGAG 198

•
•
•
•

OM of: US-09-932-678-2 to: Issued_Patents_NA.* out_format: pls

Date: Jul 30, 2002 7:40 AM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 CompuGen Ltd.

Command line parameters:

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-Db=Issued_Patents_NA -Opm=fastap -SufFix=oli.rni -GAPOP=4.500
-GAPEXT=0.050 -MINMATCH=0.100 -LOOPECL=0.000 -LOOPEXT=0.000
-GAPOP=4.500 -GAPEXT=0.050 -GAPOP=60.000 -GAPEXT=60.000
-GAPOP=6.000 -GAPEXT=7.000 -GAPOP=60.000 -GAPEXT=60.000
-DELOP=6.000 -DELEXT=7.000 -START=1 -MATRIX=oligo
-FRAM=human40.cdt -LIST=45 -DOCALIGN=200 -THR_SCORE=quality
-THR_MIN=1 -ALIGN=15 -MODE=LOCAL -OUTFMT=pls -NORM=ext
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-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -NO_XLPPY -WAIT -THREADS 1

Search information block:

Query: US-09-932-678-2

Query length: 651

Database: Issued_Patents_NA.*

Database sequence: 38353

Database length: 122816752

Search time (Sec): 52.360000

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WARN: XGAPEXT and YGAPEXT must be equal. Assuming YGAPEXT=XGAPEXT=60.000

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/cgn2_6/ptodata/1/ina/6A.COMB.seq:US-08-617-454-1 +		8.00	107.72	259.30	3901	1
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/cgn2_6/ptodata/1/ina/6A.COMB.seq:US-08-720-229-96 +		7.00	106.51	302.85	386	1
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seq_documentation_block:

Sequence 202, Application US/08905223

Patent No. 6222023

GENERAL INFORMATION:

APPLICANT: Edwards, Jean-Baptiste D.

APPLICANT: Dueller, Aymoric

APPLICANT: Lacroix, Bruno

TITLE OF INVENTION: 5' ESTS FOR SECRETED PROTEINS

NUMBER OF SEQUENCES: 503

CORRESPONDENCE ADDRESS:

ADDRESSEE: Knobbe, Martens, Olson & Bear

STREET: 501 West Broadway

CITY: San Diego

STATE: California

COUNTRY: USA

ZIP: 92101-4505

COMPUTER READABLE FORM:

SEQUENCE TYPE: Platy disk

COMPUTER: IBM pc compatible

OPERATING SYSTEM: Windows

SOFTWARE: word

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/905.223

FILING DATE:

CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:

NAME: Isgel, Sen, Ned A.

REGISTRATION NUMBER: 29,655

REFERENCE/DOCKET NUMBER:

TELECOMMUNICATION INFORMATION:	
TELEPHONE: (619) 235-8550	
TELEFAX: (619) 235-0176	
INFORMATION FOR SEQ ID NO: 202:	
SEQUENCE CHARACTERISTICS:	
LENGTH: 309 base pairs	
TYPE: NUCLEIC ACID	
STRANDEDNESS: DOUBLE	
TOPOLOGY: LINEAR	
MOLECULE TYPE: cDNA	
ORGANISM: Homo Sapiens	
ISOR TYPE: Br 31n	
FEATURE:	
NAME/KEY: other	
LOCATION: 105..252	
IDENTIFICATION METHOD: blastn	
OTHER INFORMATION: Identity 94	
OTHER INFORMATION: region 136..283	
OTHER INFORMATION: id HS046355	
OTHER INFORMATION: est	
FEATURE:	
NAME/KEY: other	
LOCATION: 227..276	
IDENTIFICATION METHOD: blastn	
OTHER INFORMATION: Identity 94	

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seq_name: /cruz/production/1/ina/6b_c0mb.seq;US 09 500 569 54
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seq_documentation_block:
: Sequence 9, Application US/09500569
: Patent No. 6329204
: GENERAL INFORMATION:
: APPLICANT: Cahoon, Rebecca R.
: APPLICANT: Katsuki, Antoni
: APPLICANT: Shen, Jennie
: TITLE OF INVENTION: Plant Cellule acid 3-O Methyltransferase Homologs
: FILE REFERENCE: B1327 US NA
: CURRENT APPLICATION NUMBER: US/09/500,569
: CURRENT FILING DATE: 2000-02-09
: EARLIER APPLICATION NUMBER: 60/119,587
: EARLIER FILING DATE: 1999-February-10
: NUMBER OF SEQ ID NOS: 28
: SOFTWARE: Microsoft Office 97
: SEQ ID NO 9
: LENGTH: 1342
: TYPE: DNA
: ORGANISM: Oryza sativa
US-09-500-569-9

alignment_scores:
Quality: 8.00 Length: 8
Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-932-678-2 x US-09-500-569-9/rev ..
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14 AlAAlaAlaSerSerAlaVal 21
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235 GCCCGCCGCGCGTCGCGCGGTC 212

seq_name: /cqn2_6/ptodata/1/ina/6A_COMB.seq:US-08-815-809-2

seq_documentation_block:
: Sequence 2, Application US/08815609
: Patent No. 6004777
: GENERAL INFORMATION:
: APPLICANT: TARTAGLIA, James
: APPLICANT: GOEBEL, Scott J.
: APPLICANT: COX, William I.
: APPLICANT: GETTIG, Russell R.
: APPLICANT: PINCUS, Steven E.
: APPLICANT: PAOLETTI, Enzo
: APPLICANT: JACOBS, Hettiam L.
: TITLE OF INVENTION: VECTORS HAVING ENHANCED EXPRESSION, AND METHODS OF
: TITLE OF INVENTION: MAKING AND USES THEREOF
: FILE REFERENCE: 454310-1010
: CURRENT APPLICATION NUMBER: US/08/815,809
: CURRENT FILING DATE: 1997-03-12
: NUMBER OF SEQ ID NOS: 23
: SOFTWARE: PatentIn Ver. 2.0
: SEQ ID NO 2
: LENGTH: 2844
: TYPE: DNA
: ORGANISM: Vaccinia virus
US-08-815-809-2

alignment_scores:
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Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

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Align seq 1/1 to: US-08-815-809-2 from: 1 to: 2844

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541 AtGsetThAlaGlyAlaSer 548
111111111111111111111111
2564 GATCAACGCGCTGCTGGGATAGC 2607

seq_name: /cqn2_6/ptodata/1/ina/5B_COMB.seq:US-08-816-155B-4

seq_documentation_block:
: Sequence 4, Application US/08816155B
: Patent No. 5990091
: GENERAL INFORMATION:
: APPLICANT: TARTAGLIA, James
: APPLICANT: COX, William I.
: APPLICANT: GETTIG, Russell R.
: APPLICANT: MARTINEZ, Hector
: APPLICANT: PAOLETTI, Enzo
: APPLICANT: PINCUS, Steven E.
: TITLE OF INVENTION: VECTORS HAVING ENHANCED EXPRESSION, AND
: TITLE OF INVENTION: METHODS OF MAKING AND USES THEREOF
: NUMBER OF SEQUENCES: 48
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: FROMMER LAWRENCE & HAUG LLP
: STREET: 745 FIFTH AVENUE
: CITY: NEW YORK
: STATE: NEW YORK
: COUNTRY: USA
: ZIP: 10151
: COMPUTER READABLE FORM:
: MEDIUM TYPE: floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.40
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/816,155B
: FILING DATE: 12-MAR-1997
: CLASSIFICATION: 514
: ATTORNEY/AGENT INFORMATION:
: NAME: KOMALSKI, THOMAS J.
: REGISTRATION NUMBER: 42,147
: REFERENCE/DOCKET NUMBER: 454410-2990
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 212-588-0800
: TELEFAX: 212-588-0500
: INFORMATION FOR SEQ ID NO: 4:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 2856 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
US-08-816-155B-4

alignment_scores:
Quality: 8.00 Length: 8
Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-932-678-2 x US-08-816-155B-4 ..
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111111111111111111111111
2594 GATCAACGCGCTGCTGGGATAGC 2619

seq_name: /cqn2_6/ptodata/1/ina/6A_COMB.seq:US-09-079-587-4

seq_documentation_block:
: Sequence 4, Application US/09079587
: Patent No. 6130066
: GENERAL INFORMATION:

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: CLASSIFICATION: 800
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 08/012,688
: FILING DATE: 03-FEB-1993
: ATTORNEY/AGENT INFORMATION:
: NAME: Lavin J., Lawrence M.
: REGISTRATION NUMBER: 30,768
: REFERENCE/DOCKET NUMBER: 38-21(13583)A
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (314)537-6670
: TELEFAX: (314)537-6047
: INFORMATION FOR SEQ ID NO: 1:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 3901 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: US-08-617-454-1

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  Ratio: 1.000      Gaps: 0
  Percent Similarity: 100.000  Percent Identity: 100.000

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seq_name: /cgn2/6/plodata/1/ina/PCTUS_COMB.seq:PCT-US94-01144-1
seq_documentation_block:
: Sequence 1, Application PC/TUS9401144
: GENERAL INFORMATION:
: APPLICANT:
: TITLE OF INVENTION: Plants Resistant to Infection by FLRV
: NUMBER OF SEQUENCES: 5
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: PCT/US94/01144
: FILING DATE: 01-FEB-1994
: CLASSIFICATION:
: INFORMATION FOR SEQ ID NO: 1:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 3901 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: cDNA (genomic)
: PCT-US94-01144-1

alignment_scores:
  Quality: 8.00      Length: 8
  Ratio: 1.000      Gaps: 0
  Percent Similarity: 100.000  Percent Identity: 100.000

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Align seq 1/1  to: PCT-US94-01144-1  from: 1  to: 3901
582 GIUASPMetSerAlaGluGluLeu 589
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|||||
2293 GAGGATATCAGCCGACAGACAGCTG 2316

seq_name: /cgn2/6/plodata/1/ina/6A_COMB.seq:US-08-815-809-3
seq_documentation_block:
: Sequence 3, Application US/08815809
: Patent No. 6004777
: GENERAL INFORMATION:
: APPLICANT: TAKAGIYA, James
: APPLICANT: GORHEL, Scott J.
: APPLICANT: COX, William L.
: APPLICANT: GETTIG, Russell R.
: APPLICANT: PINCUS, Steven E.
: APPLICANT: PROFFIT, Enzo
: APPLICANT: JACOBS, Bettam L.
: TITLE OF INVENTION: VECTORS HAVING ENHANCED EXPRESSION, AND METHODS OF
: TITLE OF INVENTION: MAKING AND USES THEREOF
: PTE REFERENCE: 454310-3010
: CURRENT APPLICATION NUMBER: US/08/815,809
: CURRENT FILING DATE: 1997-03-12
: NUMBER OF SEQ ID NOS: 23
: SOFTWARE: PatentIn Ver. 2.0
: SEQ ID NO: 3
: LENGTH: 6628
: TYPE: DNA
: ORGANISM: Vaccinia virus
: US-08-815-809-3

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  Quality: 8.00      Length: 8
  Ratio: 1.000      Gaps: 0
  Percent Similarity: 100.000  Percent Identity: 100.000

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  US-09-932-678-2 x US-08-815-809-3/rev  ..

Align seq 1/1  to reverse of: US-08-815-809-3  from: 1  to: 6628
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seq_name: /cgn2/6/plodata/1/ina/5H_COMB.seq:US-08-816-155B-5
seq_documentation_block:
: Sequence 5, Application US/08816155B
: Patent No. 5990991
: GENERAL INFORMATION:
: APPLICANT: TAKAGIYA, JAMES
: APPLICANT: COX, WILLIAM L.
: APPLICANT: GETTIG, RUSSELL R.
: APPLICANT: MARINEZ, HECTOR
: APPLICANT: PALETTI, ENZO
: APPLICANT: PINCUS, STEVEN E.
: TITLE OF INVENTION: VECTORS HAVING ENHANCED EXPRESSION, AND
: TITLE OF INVENTION: METHODS OF MAKING AND USES THEREOF
: NUMBER OF SEQUENCES: 48
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: FROMMER, LAWRENCE & HANG LLP
: STREET: 745 FIFTH AVENUE
: CITY: NEW YORK
: STATE: NEW YORK
: COUNTRY: USA
: ZIP: 10151
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/816,155B
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TEL: 011-26111111 FAX: 011-26111111

1. *Pharyngodon* *liberal*

DS-108-244-788A-4(1)

alignment_scores:
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Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

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39 AGTCGACCGCGGGGGGAT 19

seq_name: /cyn2_6/prodata/1/ina/5A_COMB.seq:US-08-049-473-19

seq_documentation_block:

: Sequence 19, Application US/08049473
: Patent No. 5386021

: GENERAL INFORMATION:

: APPLICANT: Moss, Joel

: APPLICANT: Mishima, Koichi

: APPLICANT: Nishimura, Maria

: APPLICANT: Tsuchiya, Mikako

: TITLE OF INVENTION: A MAMMALIAN GUANIN NUCLEOTIDE BINDING

: NUMBER OF SEQUENCES: 34

: CORRESPONDENCE ADDRESS:

: ADDRESSEE: KNOBBE, MARTENS, OLSON AND BEAR

: STREET: 620 NEWPORT CENTER DRIVE SIXTEENTH FLOOR

: CITY: NEWPORT BEACH

: STATE: CA

: COUNTRY: USA

: ZIP: 92660

: COMPUTER READABLE FORM:

: MEDIUM TYPE: Floppy disk

: COMPUTER: IBM PC compatible

: SOFTWARE: Patent Release #1.0, Version #1.25

: CURRENT APPLICATION DATA:

: APPLICATION NUMBER: US/08/049,473

: FILING DATE: 19930419

: CLASSIFICATION: 436

: ATTORNEY/AGENT INFORMATION:

: NAME: Fuller, Michael L.

: REGISTRATION NUMBER: 36,516

: REFERENCE/DOCKET NUMBER: NIH050,001CP1

: TELECOMMUNICATION INFORMATION:

: TELEPHONE: 619-235-8550

: TELEFAX: 619-235-0176

: INFORMATION FOR SEQ ID NO: 19:

: SEQUENCE CHARACTERISTICS:

: LENGTH: 48 base pairs

: TYPE: nucleic acid

: STRANDEDNESS: single

: TOPOLOGY: linear

: MOLECULE TYPE: cDNA

: HYPOTHETICAL: NO

: ANTI-SENSE: NO

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Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

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Align seq 1/1 to reverse of: US-08-049-473-19 from: 1 to: 48

170 SerAspSerAspAspGluAsp 176
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35 TGGACTCGGACGATGAGGAT 19

LOCUS AL597171 550 bp mRNA linear EST 14-AUG-2001
 DEFINITION DKFZP313K0411.1 313 (synonym: hlec2) Homo sapiens cDNA clone
 ACCESSION AL597171
 VERSION AL597171.1 GI:15154983
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 REFERENCE 1 (bases 1 to 550)
 Koehler, K., Beyer, A., Mewes, W., Weil, H. and Wiemann, S.
 EST (Koehler, K., Beyer, A., Mewes, W., Weil, H. and Wiemann, S.)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Koehler K
 MFS
 Am Klopferspitz 18a D-82152 Martinsried, Germany
 This is the 5' sequence of the clone insert
 clone from S. Wiemann, Molecular Genome Analysis, German Cancer
 Research Center (DKFZ), Email: s.wiemann@dkfz-heidelberg.de;
 sequenced by BMFZ (Biomedical Research Center at the Charité,
 Berlin/Germany) within the cDNA sequencing consortium of the German
 Genome Project.
 No sl sequence available.
 This clone (DKFZP313K0411) is available at the RZPD in Berlin.
 Please contact the RZPD: Rossourcezentrum, Heubnerweg 6, 14059
 Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.
 Location/Qualifiers
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 /db_xref="taxon:9606"
 /clone="DKFZP313K0411"
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 /dev_stage="adult"
 /lab_host="DH10B"
 /note="Vector: pTRipleX2; site_1: S11A; site_2: S11B;
 cDNA-collection"
 BASE COUNT 140 a 116 c 140 g 153 t 1 others
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 Ratio: 1.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000
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 17 rsetSerAlaValIlyslsleuGlyAlaSerArthrGlyIleSerAsnM 34
 98 GTCCTGTGCACTTAAGACGTCGGCGCGCTCGAGGAGCTGCTTCAATA 147
 34 eTAAGAlaLeuGluAsnAspPheAsnSerProProArthrLysThrVal 50
 148 TCGCGTCATTAGACAAATGACTTTTCAATTTCCGCCCAAGAAAAGCTTT 197
 51 ArapheGlyIlyThrValThrGluValLeuLeuLysTyrIlyslsGly 67
 198 CGGTTTGTGCACTGTGACAGAGCTCTGCTGAAGTACAAAGACGCTGA 247
 67 uThAsAspPheGluLeuLeuLysAsnGlnLeuLeuAspProAspIleL 84
 248 AACCAATGACTTTGAGTTGTTGCAAGAACCACTGTTAGATVTCACACATA 297
 84 YSASPASGlnIleIleAsnTrpLeuLeuGluPheArgSerSerIleMet 100
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258 AGCATGACATGATGACATGCTGTGTAATTCGATTCTTATGATG 447
 101 TgtLeuThrIlyAspPheGluGlnLeuIleSerIleIleLeuArLeuP 117
 348 TACTGTGCAAAAGACATTCAGCAACTATATGATATATATAGATTTC 307
 117 oTTPLeuAsnArgSerGluThrValValGluGluGlyTyrLeuAlaPheGlu 144
 398 TTGCTTAAATGACAGTAAACACTAGTCAGAACGATTTGCTTTCTG 447
 134 lYAsnLeuValSerAlaGlnThrValPheLeuArgProCysLeuSerMet 150
 448 GATATCTGTATGACACACACAGCTTTCTTCACACGCTGTCTCATGATG 497
 151 lTleAlaSerHisPheValProProArthrValIleIleLysGluIlyAsp 167
 498 ATGCTTCCCATTTGTGCTTCCCGCATGATATAGAGAGATGATG 547
 167 1 167
 548 A 548
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 seq_documentation_block:
 LOCUS AA213789 545 bp mRNA linear EST 14-AUG-1997
 DEFINITION 7791a11.st NC1_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:680056.4.
 mRNA sequence.
 ACCESSION AA213789
 VERSION AA213789.1 GI:1812416
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 REFERENCE 1 (bases 1 to 545)
 NC1_CGAP <http://www.ncbi.nlm.nih.gov/nc1cgap>.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 JOURNAL Unpublished (1997)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: rga@bbs-rtm1.nih.gov
 This clone is available royalty-free through INM; contact the
 IMAG Consortium (info@image.lln.gov) for further information.
 Possible reversed clone; polyT not found
 Insert length: 1076 Std Error: 0.00
 Seq primer: 41m13 fwd, 5' from Amersham
 High quality sequence stop: 457.
 Location/Qualifiers
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 1..545
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 /db_xref="taxon:9606"
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 /issue_type="germline center B cell"
 /lab_host="DH10B"
 /note="Vector: pTZ19-Pac (Pharmacia) with a modified
 polylinker; site_1: Not I; site_2: Eco RI; 1st strand cDNA
 was prepared from human tonsillar cells cultured for
 germinal center B cells by flow sorting (Ch20c, 10c+),
 provided by Dr. Louis M. Staudt (NCI), Dr. David Allman
 (NCI) and Dr. Gerald Mark (CBER). cDNA synthesis was
 primed with a Not I - oligo(dT) primer
 5'-TGTACCAATCTGAATGAGCGCGGCGCTCATTTTCTTTTCTTTT
 1. Double-stranded cDNA was ligated to Eco RI adaptors
 (Pharmacia), digested with Not I and cloned into the Not I
 and Eco RI sites of the modified pTZ19 vector. Library
 went through one round of normalization, and was
 constructed by Benito Soares and M. Fatima Bonaldo.
 BASE COUNT 150 a 102 c 121 g 161 t 1 others

seq_documentation_block:
LOCUS BG167061 990 bp mRNA linear EST 06-FEB-2001
DEFINITION 6023447401 NIH_MGC_89 Homo sapiens cDNA clone IMAGE:445009 5',
mRNA sequence.
ACCESSION BG167061
VERSION BG167061.1 GI:12673764
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE 1 (bases 1 to 990)
AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: rsb@biml.nih.gov
Tissue procurement: ATCC
cDNA library preparation: Life Technologies, Inc.
cDNA library arrayed by: The T.M.A.G.E. Consortium (LMN)
DNA sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the T.M.A.G.E. Consortium/LMN at:
<http://image.llnl.gov>
Plate: LAM10247 row: 0 column: 10
High quality sequence stop: 665.
FEATURES
source location/Qualifiers
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/db_xref="taxon:9606"
/clone_lib="NIH_MGC_89"
/tissue_type="hypopharynx, cell line"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: Kidney; Vector: pCMV-SPOR16; Site: 1; Not:
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full-length clones and constructed by Life Technologies.
Note: This is a NIH-MGC library."
BASE COUNT 304 a 275 c 219 g 192 t
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Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000
alignment_block:
US-09-932-678-2 x BG167061 ..
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108 GCGCTAGAGATTGCGCGCCCTAGGCGTTACTTTTCTGCTCAATCAC 157
517 fasnlstytyglaleuvalphecytytrhrlelelelelarsasnasna 534
|||||
158 AATTAATACAGCTGCTCTTCTGCTAGACCATTTGAGAGACATC 207
534 tgcloleleuprovallelarssethralaslyclaspservalgla 550
|||||
208 GGCACATGCTGCCAGTCATTAAGAGTACCGCTGAGAGACATCAGTCAG 257
551 llecystrhasnproleuaspphrphepheprophasprocyasvlla 567
|||||
258 ATCTGACAAACCGCGGACACCTCTCTCCCTTTGATCCCTGTGTGCT 307
567 ulysarsetlyslsphenleasprolelelyglvaltrpglnaspm 584
|||||
308 CAAGAGCTCAAGCAATTCATTCATTCATTATGACGCTGTGGACACA 357

584 cctsetalactactlucenuglnlupheylslyprometlylsasplle 600
|||||
358 TAGAGTCAAGAGCTACAGAGTTTAAAGAAACATCAAAAAGGATA 407
601 valgluaspliaspaspaspheleuylslyglvalprogluasnas 617
|||||
408 GTCGAGATCAAGATGATCATTCTTGAAAGCGGAGAGTCCTGCACATTA 457
617 pthrvallllygllylethrprosetserphesapthrlhspluarqsp 634
|||||
458 TACCTGATGGATGATCAACACAGACCTGCTTTGACAGCGCATTTGCAATG 507
634 rosetsersevalylserspro 641
|||||
508 CTTCAGATAGTGTGGGCTGCCCA 530
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seq_documentation_block:
LOCUS BE093077 422 bp mRNA linear EST 12-JUN-2000
DEFINITION KC5-BF0744-260400-031 H10 BF0744 Homo sapiens cDNA, mRNA sequence.
ACCESSION BE093077
VERSION BE093077.1 GI:8483529
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE 1 (bases 1 to 422)
AUTHORS Dias Neto, E., Garcia Cortes, R., Veljovski, Alimida, S., Jimenez, M.,
Nadal, M.A., da Silva, M., Jr., Zago, M.A., Jordao, S., Costa, F.,
Goldman, G.H., Carvalho, A.F., Matsukuma, A., Paula, D.S., Simpson, J.H.,
Brunstein, A., de Oliveira, P.S., Bucher, P., Jongsomjit, C.V., O'Hare,
M.J., Soares, F., Brenhan, R.R., Reis, L.F., de Souza, S.J., and
Simpson, A.J.
TITLE Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
MEDLINE 20202663
COMMENT Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Judeq Institute for Cancer Research
Judeq Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@judeq.org.br
This sequence was derived from the FAPESP/PCR Human Cancer Genome
Project. This entry can be seen in the following DB:
(<http://www.judeq.org.br/scripts/acthtml2.pl?1> &2-405 &10744-260
400-031-BF0744-260004-26044-1)
Seq primer: puc 18 forward
High quality sequence stop: 414.
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/clone_lib="BF0744"
/seq_stage="Adult"
/notes="Organ: breast; Vector: puc18; Site: 1; Smal: Site 2;
Smal: A mini-library was made by cloning products derived
from ORESTES PCR (U.S. Letters Patent application No. 196
7716 - Judeq Institute for Cancer Research) products
into the pUC 18 vector. Reverse transcription of tissue
mRNA and cDNA amplification were performed under low
stringency conditions."
BASE COUNT 121 a 98 c 99 g 104 t
ORIGIN
alignment_scores:
Quality: 140.00 Length: 140


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DEFINITION      zv02e05.f1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:746336 5',
                  mRNA sequence.
ACCESSION       AA481295
VERSION         AA481295.1 GI:2210847
KEYWORDS        EST.
SOURCE          human.
ORGANISM        Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE       1 (bases 1 to 380)
                 NCI_CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
                 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
                 Tumor Gene Index
                 Unpublished (1997)
JOURNAL         Contact: Robert Strausberg, Ph.D.
                 Email: cgapsb@mail.nih.gov
                 Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
                 Ph.D., Gerald Marti, M.D.
                 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
                 Ronaldo, Ph.D.
                 cDNA Library Arrayed by: Greg Lennon, Ph.D.
                 DNA Sequencing by: Washington University Genome Sequencing Center
                 Clone distribution: NCI-CGAP clone distribution information can be
                 found through the I.M.A.G.E. Consortium/LIN. at:
                 www.bio.lnln.gov/biopr/image/image.html
                 Seq primer: -28m3 rev1 ET from Amersham
                 High quality sequence stop: 361.
FEATURES        location/Qualifiers
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                 /issue_type="germinal center B cell"
                 /lab_host="DH10B"
                 /note="Vector: pTZ19-Pac (Pharmacia) with a modified
                 polylinker. Site_1: Not 1; Site_2: Eco RI; 3rd strand cDNA
                 was prepared from human tonsillar cells enriched for
                 germinal center B cells by flow sorting (CD20+, IgD-),
                 provided by Dr. Louis M. Staudt (NCI), Dr. David Allman
                 (NCI) and Dr. Gerald Marti (CBER). cDNA synthesis was
                 primed with a Not I - oligo(dT) primer
                 15'-TGTTACCATCTGAGTGGGAGCGCCCTCATTTTCTTTTCTTTT-3'
                 1. Double-stranded cDNA was ligated to Eco RI adaptors
                 (Pharmacia), digested with Not I and cloned into the Not I
                 and Eco RI sites of the modified pTZ19 vector. Library
                 went through one round of normalization, and was
                 constructed by Bento Soares and M. Fatima Ronaldo."
BASE COUNT      105 a 90 c 76 g 109 t
ORIGIN
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  Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000
alignment_block:
  US-09-932-678-2 x AA481295/rev ..
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379 ATTGAATATGCTGAGAAACAGCTAACTGCTGCTGGACAGATTC 330
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279 rThcIuglyLeuPheAsnMetAspGluAspGluGlnThrGluHisGlu 296
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329 CAGGGAACGATGTTTATATGATGAGATGAAGATGAAGAACTGAAATGAAA 280
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296 hrlysalaglyPProGluArfLeuAspGlnMetValHisProValAlaGlu 412
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275 CAAGAGCTGGTCTCTCAAGCGCTGACCAACATAGTGGATCTGTATGCGAG 240
313 ArleuAspLleleuMetSerLeuValIleuSerTyrMetLysAspValcy 429
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229 CGCTGGACATGCTGATGCTCTTGCTTGCTCAATGACATGATCTG 180
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329 sTyValAspGlyValValAspAsnGlyLysThrLysAspLeuTyrATCA 446
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179 CTAATGATGATGTAAGCTGATATACCCCAAAACAAAGCATATATATGCG 140
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346 sPnuLeuAsnLlePheAspLysLeuLeuLeuProThrHisAlaSerCys 362
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|||||
363 HisValGlnThrPheMetPheTyrLeuGlySerPheLysGluGlyPheAl 479
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70 CAGTCACAGTTCCTTCATGCTTACCTGCTGAGTTCATCAATGGCATTCG 40
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375 aAluAlaPheLeuGluGlnHisLeuTyrLys 388
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20 AAGCCATTTTTCGAAATATCTGTGAAA 2
seq_name: gb_cst1:AA191111
seq_documentation_block:
LOCUS      AA191111 421 bp mRNA linear EST 15 JAN 1997
DEFINITION zpb6a04.f1 Stradaque HeLa cell s3 947216 Homo sapiens cDNA clone
IMAGE:627052 5' similar to WP03688.1 CPO0909.2 mRNA sequence.
ACCESSION AA191111
VERSION   AA191111.1 GI:1779805
KEYWORDS  EST.
SOURCE    human.
ORGANISM  Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE  1 (bases 1 to 421)
AUTHORS   Hillier, L., Lennon, G., Becker, M., Bonaldo, M.F., Chapple, B.,
           Chissole, S., Dietrich, N., Dubouche, T., Favello, A., Gish, W., Hawkins,
           M., Hultman, M., Kucaba, T., Lacy, M., Le, M., Le, N., Maridis, E., Moore,
           B., Morris, M., Parsons, J., Prange, C., Rifkin, B., Rohlfing, J.,
           Schellengberg, K., Soares, M.B., Tan, F., Tibbitts, M., Trevisan, E.,
           Underwood, K., Wohlmann, P., Waterston, P., Wilson, R. and Marra, M.
           Generation and analysis of 280,000 human expressed sequence tags
           Genome Res. 6 (9), 807-828 (1996)
97044478
CONTACT: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63110
Tel: 314 286 1800
Fax: 314 286 1810
Email: est.watson@wustl.edu
This clone is available royalty-free through LIN. Contact the
IMAGE Consortium (infoimage.lnln.gov) for further information.
Seq primer: -28m3 rev2 from Amersham
High quality sequence stop: 383.
location/Qualifiers
1..421
/organism="Homo sapiens"
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/lab_host="Solk (kanamycin resistant)"
/note="Vector: pBluescript SK-; Site 1: EcoRI; Site 2:
XhoI; cloned unidirectionally. Primer: clone d1, HeLa S3
epitheloid carcinoma cells grown to semi-confluent
without induction. Average insert size: 1.5 kb; 0.5 ZAP-XR
Vector. -5' adaptor sequence: 5' GAAATCGGTAATGAT 3'
adaptor sequence: 5' CTGACGTTTCTTTTCTTTTCTTTT 3'"
BASE COUNT      108 a 93 c 86 g 134 t
ORIGIN

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DEFINITION      6018631f1 NIH_MGC_17 Homo sapiens cDNA clone IMAGE:4099398 5',
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ACCESSION       AF203684
VERSION         AF203684.1 GI:11097270
KEYWORDS        EST.
SOURCE          human.
ORGANISM        Homo sapiens
                Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE       1 (bases 1 to 996)
                NIH-MGC http://mgc.nhl.nih.gov/.
AUTHORS         National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE           Unpublished (1999)
JOURNAL         Contact: Robert Strausberg, Ph.D.
                Email: cyabbs@mail.nih.gov
COMMENT         Tissue Procurement: ATCC
                cDNA Library Preparation: Ling Hong/Rubin Laboratory
                DNA Sequencing by: Incyte Genomics, Inc.
                Clone distribution: MGC clone distribution information can be
                found through the I.M.A.G.E. Consortium/LINL at: image.lnl.gov
                Plate: LLCM967 row: n column: 07
                High quality sequence stop: 684.
                Location/Qualifiers
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                        /lab_host="DH10B (phage-resistant)"
                        /note="organ: muscle; Vector: pOTB7; Site_1: EcoRI;
                        Site_2: XhoI; cDNA made by oligo-dt priming.
                        directionally cloned into EcoRI/XhoI sites using the
                        following 5' adaptor: GGCACGAG(C). Size-selected >500bp
                        for average insert size 1.8kb. Library constructed by
                        Ling Hong in the laboratory of Gerald M. Rubin (University
                        of California, Berkeley) using ZAP-cDNA synthesis kit
                        (Stratagene) and Superscript II RT (Life Technologies)."
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BASP COUNT      338 a      211 c      231 g      216 t
ORIGIN
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    Percent Similarity: 100.000      Percent Identity: 100.000
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155 GTTAAGTGGCTGGCAGATATCTTAATATACGACATTTGGGACACAAAGGC 204
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446 aPheCysAspValAlaLeuHisGlyProPheTyrSerAlaCysGlnAlaVal 463
|||||
205 ATTTCGACATGTTGCTCTCCATGACCATTTTACTACGCTGCACAGCTG 254
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463 aPheTyrThrPheValPheArhHisIleGlnLeuLeuSerGlyAsnLeu 479
|||||
255 TGTTCATACCTTGTGTTTATAGACACAGCAGCTTTTGACCGCAACCTG 304
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480 LysGluGlyLeuGlnTyrLeuGlnSerLeuAsnPhenGlnArgIleValIle 496
|||||
305 AAAGAGAGCTTTTCAGATATCTTCAGAGCTGTGAATTTTGACGATAGTGAT 354
|||||
496 tSerGlnLeuAsnProLeuLysIleCysLeuProSerValValAsnPheP 513
|||||
355 GAGCCAGCTAAATCCCTGAACATTTGCCCTGCCCTCACTGTTAACTTTT 404
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513 LeuAlaAlaIleThrAsnLysTyrGlnPheValPheCysTyrThrIleIle 529
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405 TTGCTGCAATCAACAATAAGTACAGTTCGCTTCTGCTAGACATCATCAT 444
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530 GUAATGASAsnArgGlnMetLeuProValIleArgSerThrAlaLysAl 546
|||||
455 GAGAGGAGACATCGCCGAGATCGTCCGAGCTCATTAAGAGTACGCTGGAG 504
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546 yAspSerValGlnIle 551
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505 AGACTCAGTCCAGATTC 520
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OM of: US-09-932-678-2 to: GenBank: * out_format: pts

Date: Jul 30, 2002 7:35 AM

About: Results were produced by the GenCue software, version 4.5.
Copyright (c) 1993-2000 CompuGen Ltd.

Command line parameters:

-MOTIF: frame, p2n, model -DEV: x1h
-V: /c:/usr/local/spool/US09932678/runtal_29072002_155155_19010/app_query.fasta.1.717
-Dh: GenBank -QFMT: fastap -SUFFIX: oli.rge -GAP: 4.500
-GAPEXT: 0.050 -MINMATCH: 0.100 -LOOPT: 0.000 -LOPEXT: 0.000
-GAP: 4.500 -GAPEXT: 0.050 -XGAP: 60.000 -XGAPEXT: 60.000
-DEGAP: 6.000 -DEGAPEXT: 7.000 -YGAP: 60.000 -YGAPEXT: 60.000
-DEGAP: 6.000 -DEGAPEXT: 7.000 -START: 1 -MATRIX: oli1ao
-TRANS: human40.cdt -LIST: 45 -LOCAL: 200 -THR_SCORE: quality
-THR_MIN: 1 -ALIGN: 15 -MODE: local -OUTFMT: pts -NORM: ext
-HEAPSIZE: 500 -MINLEN: 0 -MAXLEN: 200000000
-USER: US09932678 -MOTIF: 1.4676 -NCP: 6 -ICPU: 3 -LONG: 105
-DEV_TIME: 0.120 -WARN_TIME: 0.30 -NO_XLPEXT -WAIT -THREADS: 1

Search information block:

Query: US-09-932-678-2

Query length: 651

Database: GenBank: *

Database sequences: 1797656

Search time (sec): 187333701

Search time (sec): 2262.420000

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WARN: XGAPEXT and YGAPEXT must be equal. Assuming YGAPEXT=XGAPEXT=60.000

Score list:

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gb_pt:HS272050	550.00	1071.41	0.0	3756	AF2272050 Homo sapiens RNA3 mr
gb_pt:AX036047	463.00	9014.34	0.0	2040	AX036047 Sequence 1 from Patent
gb_pt:AK055742	94.00	1797.64	9.1e-92	2493	AK055742 Homo sapiens cDNA FLJ
gb_pt:AX274907	70.00	1331.94	7.9e-66	1461	AX274907 Sequence 172 from Pat
gb_pt:BC009198	68.00	1294.29	2.0e-55	1177	BC009198 Homo sapiens, Similat
gb_pt:AC004158	62.00	1145.41	2.2e-55	129473	AC004158 Homo sapiens, Similat
gb_pt:AC0092137	62.00	1144.61	2.2e-55	143900	AC0092137 Homo sapiens chromo
gb_pt:AC007615	62.00	1144.26	2.2e-55	175691	AC007615 Homo sapiens chromo
gb_pt:AC0092562	62.00	1142.82	2.7e-55	187643	AC0092562 Papio hamadryas cl
gb_pt:AC009130	62.00	1142.68	3.0e-55	191496	AC009130 Homo sapiens chromo
gb_pt:AC0090933	62.00	1142.13	3.0e-55	208008	AC0090933 Homo sapiens chromo
gb_pt:AC0097268	62.00	1140.96	3.4e-55	247331	AC0097268 Pan troglodytes cl
gb_pt:BC006441	61.00	1152.01	8.4e-56	2627	BC006441 Homo sapiens, Similat
gb_pt:AC0092375	58.00	1065.15	5.8e-51	173166	AC0092375 Homo sapiens chromo
gb_pt:AC017077	58.00	1064.92	5.9e-51	179150	AC017077 Homo sapiens chromo
gb_pt:BC001549	58.00	1064.11	6.6e-51	202904	BC001549 Human Chromosome 16
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gb_pt:AC106788	43.00	771.94	1.2e-34	170611	AC106788 Homo sapiens chromo
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gb_pt:AC022642	22.00	366.55	4.7e-12	78347	AC022642 Homo sapiens clone B
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gb_pt:AC107769	15.00	232.20	0.0001	53814	AC107769 Mus musculus clone F
gb_pt:AC006150	15.00	227.51	0.0003	10800	AC006150 Homo sapiens clone F
gb_pt:AC0092119	15.00	224.38	0.0004	171940	AC0092119 Homo sapiens chromo
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gb_pt:AC0092285	13.00	186.86	0.0040	135873	AC0092285 Homo sapiens clone
gb_pt:AC014444	13.00	184.56	0.0649	132815	AC014444 Homo sapiens clone
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gb_pt:AF143927	10.00	160.39	1.43	1140	AF143927 Marmota sibirica clon
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DEFINITION Homo sapiens RNA3, complete cds.
ACCESSION AF227156
VERSION AF227156.1 GI:7670099
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryote; Metazoa; Chordata; Gracilata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
REFERENCE
1 (bases 1 to 2068)
Moorhead, B., Greene, E.A. and Reeder, R.H.
RNA polymerase II transcription factor Rn3 is functionally
conserved between yeast and human
Proc. Natl. Acad. Sci. U.S.A. 97 (9), 4724-4729 (2000)
MEDLINE
20243763
2 (bases 1 to 2068)
Moorhead, B., Greene, E.A. and Reeder, R.H.
Direct Submission
TITLE
Submitted (20-JAN-2000) Basic Sciences, Fred Hutchinson Cancer
JOURLN Research Center, 1100 Fairview Avenue N., Seattle, WA 98109, USA

FEATURES

Source

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Ratio: 651.00 Length: 651
Percent Similarity: 100.000 Percent Identity: 100.000
alignmet: block:
US-09-932-678-2 x AF227156
US-09-932-678-2 to: AF227156 from: 1 to: 2068

1..2068
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1919 TACCGATGATGGATACACAGAGCTCTTGCACCGCATTTCCGAAGTC 1968
634 rosergersvalglyserproprovalleutymelglnproserpro 650
1969 CTTCAAGTAGAGTGAGCTCCACCCGCTGTGTACATGCACCCAGCTCC 2018
651 leu 651
114
2019 CTC 2021
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seq_documentation_block:
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DEFINITION Homo sapiens mRNA for transcription initiation factor 1A protein
(TIF-1A gene).
ACCESSION AJ272050
VERSION AJ272050.1 GI:10046713
KEYWORDS TIF-1A; transcription initiation factor.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 3756)
Bodem J., Hoffmann-Rohrer U., Koss W., Deltus H., Vingron M. and
Grunml L.
TITLE Cloning and functional characterization of transcription initiation
factor TIF-1A, a growth-dependent regulator of ribosomal RNA
synthesis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 3756)
Bodem J.J.
TITLE Direct Submission
JOURNAL Submitted (07-FEB-2000) Bodem J.J., Molekulare Biologie der Zelle
11, Deutsches Krebsforschungszentrum, INF.280, 69120 Heidelberg,
GERMANY
FEATURES
source location/Qualifiers
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/evidence="experimental
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US-09-932-678-2 x HSA272050
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|||||
17 rSerSerAlaValIlystLengLYAlaSerArgThrGlyIleSeAsuM 44
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73 GCGCTCGCACTTAAGAAC TGGCGCGCGAGAGAGTGGAGATTCAAAA 122
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34 eValArgAlaLeuGlnAsnAspPheAsnSerProProAlaLysThrVal 50
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123 TCGCGCATTAAGAAAGACITTTCAATTCTCCGCCAAGAAAAATGCT 172
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51 AcgPheGlyGlyPheValThrGluValLeuLeuLysTyrLysGlyGly 67
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173 CGCTTTGGTGAACTGTGACAGAGAGCTTGTGAAGTAAAGAAAGAGTGA 222
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67 uThrAsnAspPheGlyLeuLeuLysAsnGlnLeuLeuAspProAspIle 84
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22 AACCAATAGCTTGGAGTGTTCAGAACAGCAGCTGTACATCCAGCAATA 272
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84 yAspAspSptIleIleAsnTrieuLeuGlnPheArgSerThrMet 100
|||||
27 AACATGACCAATGACATGACATGAGCTGTAGAAATTCGTTCTTACAG 322
|||||
161 TTTCTCThrLysAspPheGlnGlnGlnLeuSerIleLeuArgLeuP 117
|||||
323 TACTTGACAAAGACTTTCAGCACTTATACATTAATTAAGATGCT 372
|||||
117 CTPLGLeuAsnArgArgGlnThrValValGlnGlnTyrLeuAlaPheLeu 134
|||||
373 TCGCTTGAATACAGATGACAGCAGTACTGACAGCATATTGCTTTCTG 422
|||||
144 IAsnLeuValIserAlaGlnThrValPheLeuArgProLysLeuSerMet 150
|||||
423 GAAATCTTGAATGACAGAGCTGTTCGACACCGCTGTCCAGCATG 472
|||||
151 IleAlaSerHisPheValProProArgValIleLeuLysGlnLysPheVal 167
|||||
473 ATGCTTCCCATTTGTGCTCTCCGAGAGATATTAAAGAAAGAGATGT 522
|||||
167 IAspValIserAspSerAspAspGlnAspAsnLeuProAlaAsnProAla 184
|||||
523 AATGCTTTGCAATTTGATGATGATGATGATGATGATGATGATGATGAT 572
|||||
184 SThrTrpGlyHisArgAlaLeuGlnIleIleValaArgTyrValProSerThr 200
|||||
573 ACACAGTGTACACAGAGCTTGCAGAAATATAGCAAGATATATACCATGACA 622
|||||
201 PCTTPPheLeuMetProIleLeuValGlnLysPheProThrPheValArg 217
|||||
623 CTGTGTTTGTGATGCAATATCTGTGGAAAAATTTTCAATTGTTGAAA 672
|||||
217 SGTGTGuaGlnThrGlnGlnCysTyrValHisAsnLeuGuaArgIleSerV 234
|||||
673 ATCAAGACAAACAGCTGAATGATGATGATGATGATGATGATGATGATGAT 722
|||||
251 LeuLeuLysLysAspValAsnAlaSerArgGlnGlyIleGlnAspAlaArg 267
|||||
773 CTACTAAGTGGAGTGAATGATGATGATGATGATGATGATGATGATGAT 822
|||||
267 uThrAlaIleGlnThrCysGlyGlyThrAspSerThrGlnGlyLeuP 284
|||||


```

234 aATTpRhpEOTrLeuArgHISGluIleuGluLeuIleGluys 250
|||||
381 TATATTTCCACCTTGAGGATGAAATCTGACCTATTATTGAAAA 430
251 LeuLeuLysLeuAspValAsnAlaSerArgGlnGlyIleGluAspAlaG 267
|||||
431 CTAGCTCAAGCTGATGTGATGATCCGGCAGGATTGAAATGCTGA 480
267 ucUuTrAla 270
|||||
481 AGAACAACA 490

seq_name: qb-pr:BC009198
seq_documentation_block:
LOCUS BC009198 1177 bp mRNA linear PRI 12-JUN-2001
DEFINITION Homo sapiens. Similar to RNA polymerase I transcription factor.
ACCESSION BC009198
VERSION BC009198.1 GI:14327947
KEYWORDS MGC.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 1177)
AUTHORS Strausberg, R.
TITLE Direct Submission
JOURNAL Submitted (06-JUN-2001) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
REMARK NIH-MGC Project URL: http://mgc.ncl.nih.gov
COMMENT Contact: MGC help desk
Email: mgc@ncl.nih.gov
Tissue Procurement: Louis M. Staudt, M.D., Ph.D.
CDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Institute for Systems Biology
http://www.systemsbio.org
contact: amadan@systemsbio.org
Anup Madan, Rachel Dickhoff, Jessica Fahey, Stephanie Ford, Julia
Greene, Mark Kelleman and Anuradha Madan

Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LBL at: http://image.llnl.gov
Series: IMAGE Plate: 24 Row: K Column: 14
This clone was selected for full length sequencing because it
passed the following selection criteria: matched mRNA ql: 335302.

FEATURES
Source
1..1177
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="MGC:15121 IMAGE:3678732"
/tissue_type="Lymph, Burkitt lymphoma"
/clone_id="NH-MGC_8"
/lab_host="DH10B-R"
/role="Vector: pOTR7"
size: 636
CDS
/codon_start=1
/product="Similar to RNA polymerase I transcription factor
RNK3"
/protein_id="AAH09198.1"
/db_xref="GI:14327948"
/translation="MRALENDPFPNSPPKTVRPGSTVEVLAKTKKGFENPELLKNO
LIDPIDKDDIIMLEFRSSVMILKDEQLISILRLPWLINSQIVVEVLAISN
LVA"
BASE COUNT 339 a 222 c 253 g 363 t
ORIGIN
alignment_scores:
Quality: 68.00 Length: 68

```

```

Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000
alignment_block:
US-09-932-678-2 x BC009198
Align seq 1/1 to: BC009198 from: 1 to: 1177
31 TleSerAsnMetArgAlaLeuGluIleuAsnAspPheAsnSerProGat 47
|||||
307 ATTCAATATATGCTGTCATTAAGAGATGATTTTTCATATCTCCCTCAAG 156
47 qLysThrValArgPheGlyGlyThrValThrGluValLeuLeuLys 64
|||||
357 AAAAAGCTGTTCGTTGGTGTGCAAGCTGTCAAGAGCTTGCTGAATATA 406
|||||
64 yslsGlyGluThrAsnAspPheGluLeuLeuLysAsnGluLeuGasp 80
|||||
407 AAAAGGCTGAACCAATGATTTGATTTGCAAGAACGCTGTAGAT 456
|||||
61 ProAspIleLysAspAspGlnIleLeuAsnTrpLeuGluPheArg 97
|||||
457 CCAAGCATTAAGATGACCCACATCATCACTGCTCTAGAAATTCGTTT 506
|||||
97 rSer 98
|||||
507 TCTT 510

seq_name: qb-bc:AC040158
seq_documentation_block:
LOCUS AC040158 129473 bp DNA linear HTG 03-SEP-2000
DEFINITION Homo sapiens chromosome 16 clone G1A-13384, WORKING DRAFT SEQUENCE.
14 unordered pieces.
ACCESSION AC040158
VERSION AC040158.4 GI:5965546
KEYWORDS HTG; HTGS_PHSB1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 129473)
AUTHORS DOE Joint Genome Institute.
TITLE Sequencing of Human Chromosome 16
JOURNAL Unpublished
2 (bases 1 to 129473)
DOE Joint Genome Institute.
Direct Submission
Submitted (11-APR-2000) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
On Sep 3, 2000 this sequence version replaced gi:596552.

-----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: http://www.jgi.doe.gov
-----
Project Information
Center Project Name: 0
Center clone name: C17978SKA_13384
-----
Summary Statistics
Consensus quality: 90046 bases at least Q40
Consensus quality: 99266 bases at least Q20
Estimated insert size: 104098 bases at least Q20
Estimated insert size: 126173; sum-of-contigs estimation
Quality coverage: 4.51 in Q20 bases; adarose-1p estimation
Quality coverage: 4.4 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 14 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.

```


* 101721 122163: contig of 20443 bp in length
* 122164 122263: gap of unknown length
* 122264 143900: contig of 21637 bp in length.

FEATURES
Source
1. 143900
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="16"
/clone="RP11-344H15"
/clone_lib="RPC1 human BAC library 11"

BASE COUNT 47791 a 34128 c 34320 g 35886 t 1775 others
ORIGIN

alignment_scores:

Quality: 62.00 Length: 62
Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-932-678-2 x AC092137/rev ..

Align seq 1/1 to reverse of: AC092137 from: 1 to: 143900

421 ThrVallySerCysLeuAspLeuValAsnTrpLeuHisIleTyrLe 437
|||||
66649 ACTGTAAATCATGCTAGATCTTTGGTTAACTGCTGCACATATACCT 66600

437 uasAsnGlnAspSerGlyThrLysAlaPheCysAspValAlaLeuHisG 454
|||||
66599 TATATACAGAGATTGGGAAACAGCATTCGCGATGTTGCTCTCCATG 66550

454 LyrPheTyrSerAlaCysGlnAlaValPheTyrThrPheValPheArg 470
|||||
66549 GACCATTTACTAGCGCTCCAGCTGTGTTACACCTTTGTTTTTACA 66500

471 HisLysGlnLeuLeuSerGlyAsnLeuLysGlnGly 482
|||||
66499 CACACACAGCTTTTGACGGCAACCTGCAAGAGAGT 66464

seq_name: qb_htg:AC007615

seq_documentation_block:

LOCUS AC007615 175691 bp DNA linear HTG 03-JUL-2001
DEFINITION Homo sapiens chromosome 16 clone RP11-528K16, WORKING DRAFT
SEQUENCE 1 ordered pieces.

ACCESSION AC007615 GI:14589428

VERSION AC007615.6 HTG: HTGS_PHASE2: HTGS_DRAFT; HTGS_ACTIVEFIN.

KEYWORDS

SOURCE human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 175691)

DOE Joint Genome Institute.
Sequencing of Human Chromosome 16

Unpublished
2 (bases 1 to 175691)

BRUCE, D., MUNDT, M., DOGGETT, N., MUNK, C., SAUNDERS, E., ROBINSON, D.,
JONES, M., BUCKINGHAM, J., CHASTEN, L., THOMPSON, S., GOODWIN, L.,
BRYANT, J., TESMER, J., MEINKE, L., LONGMORE, J., WHITE, S., TATUM, O.,
CAMPBELL, C., FAWCETT, J., MALTBIE, M., BUSSOD, M., SUTHERLAND, R.,
McMURRY, K., HAN, C. and DEAVEN, L.

Direct Submission
Submitted (20-MAY-1999) Center for Human Genome Studies, DOE Joint
Genome Institute, Los Alamos National Laboratory, MS M888, Los
Alamos, NM 87545, USA

On Jul 3, 2001 this sequence version replaced qi:13928651.

Sequence Quality Assessment:
This entry has been annotated with sequence quality
estimates computed by the Phrap assembly program.
All manually edited bases have been reduced to quality zero.

Quality levels above 40 are expected to have less than

COMMENT

TITLE
JOURNAL

COMMENT

1 error in 10,000 bp.
Base-by-base quality values are not generally visible from the
Genbank flat file format but are available as part
of this entry's ASN.1 file.

Sequence Quality Assessment:

This entry has been annotated with sequence quality
estimates computed by the Phrap assembly program.
All manually edited bases have been reduced to quality zero.

Quality levels above 40 are expected to have less than
1 error in 10,000 bp.

Base-by-base quality values are not generally visible from the
Genbank flat file format but are available as part
of this entry's ASN.1 file.

* NOTE: this is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

FEATURES

Source
1. 175691
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="16"
/clone="RP11-528K16"

BASE COUNT 42392 a 42125 c 45277 g 45496 t 1 others
ORIGIN

alignment_scores:

Quality: 62.00 Length: 62
Ratio: 1.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-932-678-2 x AC007615 ..

Align seq 1/1 to: AC007615 from: 1 to: 175691

421 ThrVallySerCysLeuAspLeuValAsnTrpLeuHisIleTyrLe 437
|||||
31007 ACTGTAAATCATGCTAGATCTTTGGTTAACTGCTGCACATATACCT 31056

437 uasAsnGlnAspSerGlyThrLysAlaPheCysAspValAlaLeuHisG 454
|||||
31057 TATATACAGAGATTGGGAAACAGCATTCGCGATGTTGCTCTCCATG 31106

454 LyrPheTyrSerAlaCysGlnAlaValPheTyrThrPheValPheArg 470
|||||
31107 GACCATTTACTAGCGCTCCAGCTGTGTTACACCTTTGTTTTTACA 31156

471 HisLysGlnLeuLeuSerGlyAsnLeuLysGlnGly 482
|||||
31157 CACACACAGCTTTTGACGGCAACCTGCAAGAGAGT 31192

seq_name: qb_htg:AC092562

seq_documentation_block:

LOCUS AC092562 187643 bp DNA linear HTG 18-OCT-2001
DEFINITION Papilio hamadryas clone RP41-285113, WORKING DRAFT SEQUENCE, R
unpublished pieces.

ACCESSION AC092562 GI:16256968

VERSION AC092562.3 HTG: HTGS_PHASE1: HTGS_DRAFT; HTGS_ACTIVEFIN.

KEYWORDS

SOURCE baboon.

ORGANISM Papio hamadryas
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Wall, R., Wang, S., Ward-Moore, S., Warren, R., Washington, C.,
 Wallington, S., Williams, G., Williamson, A., Mieczys, R., Wooden, S.,
 Worley, K., Wu, C., Wu, Y., Wu, Y. F., Zhou, D., Zorrilla, S., Nelson, D.,
 Weinstein, G., and Gibbs, R.
 TITLE
 JOURNAL
 Unpublished
 Direct Submission
 2 (bases 1 to 247331)
 Worley, K.C.
 AUTHOR(S)
 TITLE
 JOURNAL
 Submitted (13-Oct-2001) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 COMMENT
 Genome Center
 Center: Baylor College of Medicine
 Center code: BCM
 Web site: <http://www.hgsc.bcm.tmc.edu/>
 Contact: hgsc-help@bcm.tmc.edu
 Project information
 Center project name: ZUAV
 Center clone name: RP43-4308

Summary Statistics
 Sequencing vector: Plasmid; M7789
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.990429
 Consensus quality: 246206 bases at least Q40
 Consensus quality: 249607 bases at least Q40
 Consensus quality: 252129 bases at least Q20
 Estimated insert size: 250832; sum-of-contigs estimation
 Quality coverage: 0x in Q20 bases; agarose-gel estimation
 Quality coverage: 8.9x in Q20 bases; sum-of-contigs estimation

NOTE: Estimated insert size may differ from sequence length
 (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)
 NOTE: This sequence may represent more than one clone.
 NOTE: This is a 'working draft' sequence. It currently
 consists of 11 contigs. The true order of the pieces
 is not known and their order in this sequence record is
 arbitrary. Gaps between the contigs are represented as
 runs of N, but the exact sizes of the gaps are unknown.
 This record will be updated with the finished sequence
 as soon as it is available and the accession number will
 be preserved.

1 58134: contig of 58134 bp in length
 * 58135 58234: gap of unknown length
 * 58235 104153: contig of 45919 bp in length
 * 104154 104253: gap of unknown length
 * 104254 142622: contig of 38369 bp in length
 * 142623 142722: gap of unknown length
 * 142723 169545: contig of 26823 bp in length
 * 169546 169645: gap of unknown length
 * 169646 190172: contig of 20527 bp in length
 * 190173 203875: gap of unknown length
 * 203876 203975: contig of 13603 bp in length
 * 203976 217146: gap of unknown length
 * 217147 217246: gap of unknown length
 * 217247 232417: contig of 15171 bp in length
 * 232418 232517: gap of unknown length
 * 232518 241219: contig of 8702 bp in length
 * 241220 245193: gap of unknown length
 * 245194 245293: contig of 3874 bp in length
 * 245294 247331: gap of unknown length
 * 247332 247331: contig of 2038 bp in length.

FEATURES

Location/Qualifiers

1..247331
 /organism="Pan troglodytes"
 /db_xref="taxon:9598"

BASE COUNT 67808 a 55778 c 54999 g 67717 t 1029 others

alignment_scores:

Quality: 62.00 Length: 62
 Ratio: 1.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-932-678-2 x AC097268/rev

Align seq 1/1 to reverse of: AC097268 from: 1 to: 247331

421 ThrVallySerCysLeuAspIleuValAsuTrpAlaHisIleTyrIle 437
 |||||
 72903 ACTGTAATGATGCTTAACTTTGGTTAACTGCTGACATATATCT 72854

437 uAsnAsnGlnAspSerGlyThrIysAlaPheCysAspValAlaLeuHisG 454
 |||||
 72853 TAAATACAGAGATTCGGCAACAGGCAATCTCCAGCTTCCTCTCAATG 72804

454 LyrProPheTyrSerAlaCysGluAlaValPheTyrThrPheValPheAs 470
 |||||
 72803 GACCATTTTACTCAGCTCTGCAAGCTGTGTCTTACACCTTTGTTTAA 72754

471 HisIysClnIleuLeuSerGlyAsnIleuLysGluGly 482
 |||||
 72753 CACCAAGACCTTTTACGGCAAACTGAAAGAGAGT 72718

seq_name: nb_pr:BC006441

seq_documentation_block:

LOCUS BC006441 2627 bp mRNA linear PK1 12 JUN 2001

DEFINITION Homo sapiens, similar to RNA polymerase I transcription factor

VERSION BC006441.1 GI:13623642

KEYWORDS MGC.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE 1 (bases 1 to 2627)

AUTHORS Strausberg, R.

TITLE Direct Submission

JOURNAL Submitted (09-Apr-2001) National Institutes of Health, Mammalian

Gene Collection (MGC), Cancer Genomics Office, National Cancer

Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,

USA

NIH-MGC Project URL: <http://mgc.ncl.nih.gov>

Contact: MGC help desk

Email: mgc@ncl.nih.gov

Tissue procurement: ATCC

cDNA Library preparation: Rubin Laboratory

cDNA Library Arrayed by: J. M. A. G. E. Consortium (IUM)

DNA sequencing by: Genome Sequence Center,

BC Cancer Agency, Vancouver, BC, Canada

Info@bcsc.bc.ca

Steven Jones, Jennifer Asano, Ian Bosdet, Yaron Buttefeld,

Susanna Chan, Readman Chin, Chris Fjell, Erin Garland, Ren Gao,

Leticia Hsiao, Martin Krzyzinski, Reta Kutsche, Oliver Levy, Soo

Sen Lee, Victor Ling, Carrie Mathewson, Candace McInerney, Steven

Ness, Pawan Pandoh, Anna-Lisa Prabhu, Farvash Sabodi, Jacques

Schlein, Diane Smalins, Michael Smith, Lorraine Spence, Jeff Stolt,

Michael Theodore, Miranada Tsai, Natasia Van den Bosch, Jill Vardy,

George Yang, Scott Zuydam, Marco Zuydam.

Clone distribution: MGC clone distribution information can be found

through the I.M.A.G.E. Consortium/UMI at: <http://imgc.llnl.gov>

Series: IRL Plate: 18 Row: d Column: 24

This clone was selected for full length sequencing because it

passed the following selection criteria: Hexamer frequency ok

FEATURES

Source

1..2627
 /organism="Homo sapiens"
 /db_xref="taxon:9606"

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 30, 2002, 02:40:21 ; Search time 1685.89 Seconds
(without alignments)
16556.064 Million cell updates/sec

Title: US-09-932-678-1

Perfect score: 2068
Sequence: 1 acagagacgctgctggaac.....taacatttgaattcccat 2068

Scoring table: OLIGO_NGC
Gapop 60.0 , Gapext 60.0

Searched: 13736207 seqs, 6748477542 residues

Word size : 0

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Host-processing: Listing first 45 summaries

Database :

EST:
1: em_estba:*
2: em_esthum:*
3: em_estin:*
4: em_estcov:*
5: em_estcov:*
6: em_estpl:*
7: em_estro:*
8: em_hic:*
9: qb_estl:*
10: qb_estl2:*
11: qb_hic:*
12: qb_qss:*
13: em_qss_hum:*
14: em_qss_inv:*
15: em_qss_pln:*
16: em_qss_vrt:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	% Match	Length	DB ID	Description
1	547	26.5	550	9	AL597171 DKFZp313K
2	510	24.7	788	10	BC502871 602550430
3	500	24.2	535	9	AA213789 zrl9a11.s
4	497	24.0	1037	10	BC428305 602498983
5	467	22.6	563	9	AW408066 01-HF-HMO
6	460	22.2	733	10	BI561523 603256222
7	458	22.1	458	10	BE549643 7b40908.x
8	449	21.7	671	9	AV703279 AV703279
9	444	21.5	564	9	AW958173 EST370243
10	445	20.1	697	10	BC611364 602612725
11	389	18.8	440	9	AA811628 0b74d03.s
12	385	18.6	996	10	BF203684 601866331
13	380	18.4	380	9	AA481295 zv02e05.r
14	373	18.1	490	10	BE502966 hz81q07.x
15	375	18.0	990	10	BC167061 602344740
16	371	17.9	422	9	BE093077 RCS-BT074
17	357	17.3	479	9	AW239267 xb38b04.y

18	337	16.4	498	10	BC149493
19	312	15.1	348	9	AW015483
20	300	14.5	421	9	AA191111
21	283	13.7	507	9	AW768543
22	278	13.4	316	10	BC149669
23	273	13.2	382	9	AV649508
24	273	13.2	828	10	BI761160
25	268	13.0	504	9	A1799954
26	267	12.9	632	10	BE614096
27	264	12.8	497	10	BE221544
28	264	12.8	498	10	BE110176
29	261	12.6	736	10	BI464721
30	260	12.6	793	10	BC501864
31	257	12.4	835	10	BI601238
32	246	11.9	461	9	A1928274
33	232	11.2	370	9	BE166014
34	218	10.5	258	10	N42382
35	217	10.5	742	10	BE870692
36	214	10.3	529	9	AW68124
37	209	10.1	448	9	AA214873
38	209	10.1	727	10	BI549394
39	205	9.9	946	10	BC535886
40	203	9.8	356	10	BE832086
41	197	9.5	253	9	AA376316
42	197	9.5	278	10	BE215433
43	196	9.5	406	10	BI018122
44	196	9.5	505	10	BC532530
45	193	9.3	327	9	AW802818

ALIGNMENTS

RESULT 1
AL597171 550 bp mRNA Linear EST 14 AUG-2002
DEFINITION DKFZp313K0411.t1 313 (Synonym: b1cc2) Homo sapiens cDNA clone
LOCUS DKFZp313K0411.5, mRNA sequence.

ACCESSION AL597171 GI:15154984
VERSION AL597171.1
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 550)
AUTHORS Koehrer,K., Beyer,A., Mewes,W., Well,B. and Wiemann,S.
TITLE EST (Koehrer,K., Beyer,A., Mewes,H.W., Well,B. and Wiemann,S.)
JOURNAL Unpublished (1999)
COMMENT Contact: Koehrer K

FEATURES
source
1..550
/organism "Homo sapiens"
/db_xref="taxon:9606"
/clone="DKFZp313K0411"
/clone_lib="113 (Synonym: b1cc2)"
/dev_stage="adult"
/lab_host="DH10B"
/vec="Vector: pT7Blue2; Site_1: SmaI; Site_2: SmaI;
Berlin-Charlottenburg, GERMANY; Email: clone-trip.dn.
location/Qualifiers
1..550

BASE COUNT 140 a 116 c 140 g 153 t 1 others


```

RESULT      3
LOCUS       AA213789
DEFINITION  z91a11.s1 NC1_CGAP_GCH1 Homo sapiens cDNA clone IMAGE:683036 3',
            mRNA sequence.
ACCESSION   AA213789
VERSION      AA213789.1 GI:1812416
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE   1 (bases 1 to 535)
AUTHORS     NC1-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE       National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
JOURNAL     Unpublished (1997)
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cgap@r-mail.nih.gov
            This clone is available royalty-free through LIND; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            Possible reversed clone: polyT not found
            Insert length: 1076 Std Error: 0.00
            Seq primer: -41ml3 fwd. fr from Amersham
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        1..535
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            /tissue_type="germinal center B cell"
            /lab_host="DH10B"
            /note="Vector: pTZ19-Pac (Pharmacia) with a modified
            polylinker site_1: Not 1; site_2: Eco RI; 1st strand cDNA
            was prepared from human tonsillar cells enriched for
            germinal center B cells by flow sorting (CD20+, IgD-),
            provided by Dr. Louis M. Staudt (NCI), Dr. David Allman
            (NCI) and Dr. Gerald Marti (CBER). cDNA synthesis was
            primed with a Not I - oligo(dT) primer
            15'-GTGTACCAATCTGACAGTGGAGCGCGCTCATTTTCTTTTCTTT-3'
            1. Double-stranded cDNA was ligated to Eco RI adaptors
            (Pharmacia), digested with Not I and cloned into the Not I
            and Eco RI sites of the modified pTZ19 vector. Library
            went through one round of normalization, and was
            constructed by Hento Soares and M. Fatima Honaldo."
BASE COUNT   150 a 102 c 121 g 161 t 1 others
ORIGIN
Query Match      24.2%; Score 500; DB 9; Length 535;
Best Local Similarity 100.0%; Pred. No. 1,3e-247;
Matches 500; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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DB 127 tctgacagaagcttcgcgaagctacaaaagctgaagaactgactgactttttaa 186
0Y 293 gaacagcgtcttatacgaacataaagaatgaacacacacacacacacacacacacacac 352
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DB 187 gaacagcgtcttatacgaacataaagaatgaacacacacacacacacacacacacacac 246

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0Y 413 atgacctgaatgaatgaagctcaaacatgaatgaagaatattagctttcttcaatga 472
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DB 307 atgacctgaatgaatgaagctcaaacatgaatgaagaatattagctttcttcaatga 406
0Y 473 tcttgatcaagaagagactgtttctcagaacatgctcgaagatgctcgaagatgctcgaat 542
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DB 427 tctgacagaagcttcgcgaagctacaaaagctgaagaactgactgactttttaa 466
0Y 593 agatgaatactctcctcaaa 612
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DB 487 agatgaatactctcctcaaa 506
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DEFINITION  BG428305.1 NIH_MGC_75 Homo sapiens cDNA clone IMAGE:3412605 5',
            mRNA sequence.
ACCESSION   BG428305
VERSION      BG428305.1 GI:13334811
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE   1 (bases 1 to 1037)
AUTHORS     NIH-MGC http://mhc.ncbi.nlm.nih.gov/.
TITLE       National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL     Unpublished (1999)
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cgap@r-mail.nih.gov
            Tissue Procurement: CLONTECH Laboratories, Inc.
            cDNA Library Preparation: CLONTECH Laboratories, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LIND)
            DNA Sequencing by: Invitrogen, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LIND at:
            http://image.llnl.gov
            Plate: LIND1361 row: e column: 22
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            /lab_host="DH10B (1) Plaque-resistant"
            /note="Organ: kidney; Vector: pUNK-11B (Clontech); Site 1:
            SmaI (agccgctcctcct); Site 2: SmaI (gacattatgac); 5' and
            3' adaptors were used in cloning as follows: 5' adaptor
            sequence: 5'-GACGCGCATATGAGC-3' and 3' adaptor sequence:
            5'-ATTCTACAGCCGACGCGCGGACATG-dT(40)NN-4' (where N = A,
            C, or G and N = A, G, C, or T). Average insert size 1.65
            kb (range 0.5-4.0 kb). 15/15 colonies contained inserts
            by PCR. This library was enriched for full length clones
            and was constructed by Clontech Laboratories (Palo Alto,
            CA). Note: this is a NIH-MGC library."
BASE COUNT   292 a 218 c 263 g 264 t
ORIGIN
Query Match      24.0%; Score 497; DB 10; Length 1037;
Best Local Similarity 100.0%; Pred. No. 4.9e-246;
Matches 497; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY	382	actttgaacacattatcaqatlatatattaagaatlgccttgatlggaatgaagaatcaaacag	441
Db	301	ACTTTGACCAACTTATTCAGTATATTATTAAAGATTCGCTTGTTGATAGAACTGCAAAACAG	360
QY	442	tatttgaagaagatatttllgqctttttcttgatgaatcttqatatacaagacaagactgttttccctca	501
Db	361	TATTGGAAGAGATATTGGCTTTCTTTGGTAATCTTGTAACAGCAAGACTGTTTTCCTCA	420
QY	502	gaacatcatctcaagatgaattgctt	525
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RESULT	10				
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ACCESSION	BC611364		mRNA sequence.		
VERSION	BC611364.1	GI:14662735			
KEYWORDS	EST.				
SOURCE	human.				
ORGANISM	Homo sapiens				

REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
1 (bases 1 to 697)	NIH-MGC	http://mgc.ncl.nih.gov/	National Institutes of Health, Mammalian Gene Collection (MGC)	Unpublished (1999)
	Contact: Robert Strausberg, Ph.D.			

cDNA Library Preparation: CLONTECH Laboratories, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MCC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LNL at:
<http://image.llnl.gov>
 Plate: L10M1607 row: a column: 08
 High quality sequence: stop: 667.

FEATURES	Location/Qualifiers
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/tissue_type="adenocarcinoma"
/lab_host="DH10b (T1 phage resistant)"
/mot-"pTad: prostate; Vector: pDR-R-LIB (Clontech);
Site-1: 5'11 (ggcgcgcgcgcgcgc); Site-2: 5'11 (ggcgcattatggc
); Double-stranded cDNA was prepared from cell line RNA.
5' and 3' adaptors were used in cloning as follows: 5'
adaptor sequence: 5'-CACGGCATTATGGC-3' and 3' adaptor
sequence: 5'-ATCTACAGCGCCGAGCGGCGGCGCATG-dT(30)BN-3'
(Where B = A, C, or G and N = A, C, G, or T). Average
insert size 1.5 kb (range 0.9-4.0 kb). 1415 colonies
contained inserts by PCR. This library was enriched for
full-length clones and was constructed by Clontech
Laboratories (Palo Alto, CA). Note: this is a NIH-MGC
Library."

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Query Match 20.1%; Score 415; DB 10; Length 697;
 Best Local Similarity 99.8%; Pred. No. 1,5e-203;
 Matches 465; Conservative 0; Mismatches 1; Indels 0;
 Gaps 0

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Qy	1723	gcaaaacaccccgctggaacacctctctccctttgaatccctgctgctgaqaqaatcgaaga	1782
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Qy	1783	aattcattgactcctattatcaggtatggaqaqaatgaqtaqtaqgaqactatgcgaat	1842
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Qy	1843	tcaagaacacccatgaaabaagaacatagtgaqaqlaaqaatgaatctctgaqaatgaq	1902
Db	319	TCACAAACCGCAAGAAAGAGACATAGTGGAAAGATGAAAGATGATGACTTTTGAAAGAGG	378
Qy	1963	aaatgcacccaabaatgaatccgtaattggagatcgaacaaactccttaaacatgaattccc	1962
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Qy	1963	gaatcctctcaatcagtgctgaatcccccacccgctttgatatagaqaqaacccatccctct	2022
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Qy	2023	gaacacgaagaatttgaaqtgaatgaatgaatctgaqaatctcccat	2068
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LOCUS	AAB1162
DEFINITION	AA811628 OB44d03.s2 NC1_CGAP_GCB1 Homo sapiens CHNA clone IMAGE:1347094 .
ACCESSION	AA811628
VERSION	AA811628.1 GI:2881239
KEYWORDS	EST.
SOURCE	human.
ORGANISM	Homo sapiens

REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
1 (bases 1 to 440)		NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap . National Cancer Institute, Cancer Genome Anatomy Project (CGAP). Tumor Gene Index Unpublished (1997)		Contact: Robert Strausberg, Ph.D.

Email: csapbs-remail.1@nih.gov
Tissue procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
Ph.D., Gerald Marti, M.D.
cDNA library preparation: M. Bento Soares, Ph.D., M. Fatima
Honaldy, Ph.D.
cDNA library Arrayed by: Greg Lennon, Ph.D.
DNA sequencing by: Washington University Genome Sequencing Center
clone distribution: NCI-CCAP clone distribution information can be
found through the I.M.A.G.E. Consortium/ILNI at:
www.bio.litni.gov/bbrp/image/image.html
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Seq primer: -40m13 fwd. Et from Amersham
High quality sequence stop: 444.
Location/Qualifiers
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/lab_host="DH10B"
/mole_vector="pTZ19 Pac (thymacia) with a modified
polylinker: Site_1: Not I; Site_2: Eco RI; 1st strand cDNA

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GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 30, 2002, 03:56:28 : Search time 58.64 Seconds

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8662.519 Million cell updates/sec

Title: US-09-932-678-1

Perfect score: 2068
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Scoring table: OLIGO-NUC

Gapop 60.0 , Gapext 60.0

Searched: 383533 seqs, 122816752 residues

Word size : 0

Total number of hits satisfying chosen parameters: 767666

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

Database : issued_patents.NA.*

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- 5: /cgn2_6/ptodata/1/lna/PCITUS.COMB.seq:*
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Prod. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	19	0.9	479	4	US-08-936-165A-192
2	19	0.9	787	1	US-08-034-245-11
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4	19	0.9	37950	4	US-09-218-207-183
5	18	0.9	315	4	US-08-744-419-3
6	18	0.9	540	4	US-08-744-419-1
7	18	0.9	708	1	US-08-078-090-24
8	18	0.9	1065	1	US-08-249-554-1
9	18	0.9	1065	1	US-08-249-555A-1
10	18	0.9	1065	1	US-08-249-584-1
11	18	0.9	1065	1	US-08-734-792-1
12	18	0.9	1065	1	US-08-078-090-1
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14	18	0.9	1065	5	PCT-US95-03789-1
15	18	0.9	1446	1	US-08-596-024-5
16	18	0.9	1446	4	US-09-020-818-5
17	18	0.9	1446	4	US-08-907-740-5
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21	18	0.9	1791	5	PCT-US95-06211-7
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23	18	0.9	6873	4	US-09-131-028A-8
24	18	0.9	8430	4	US-09-131-028A-6
25	18	0.9	8430	4	US-09-131-028A-10
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27	17	0.8	27	3	US-08-468-011A-5

28	17	0.8	27	4	US-09-236-468A-5	Sequence 5, Appl
29	17	0.8	27	5	PCT-US95-07085-5	Sequence 5, Appl
30	17	0.8	60	1	US-07-670-296-19	Sequence 19, Appl
31	17	0.8	60	1	US-08-093-781-20	Sequence 20, Appl
32	17	0.8	159	1	US-08-485-455D-78	Sequence 78, Appl
33	17	0.8	159	2	US-08-482-130C-78	Sequence 78, Appl
34	17	0.8	159	4	US-08-484-211C-78	Sequence 78, Appl
35	17	0.8	159	4	US-08-906-769-78	Sequence 78, Appl
36	17	0.8	159	3	US-08-906-616-78	Sequence 78, Appl
37	17	0.8	159	4	US-08-817-795-78	Sequence 78, Appl
38	17	0.8	159	4	US-08-485-443B-78	Sequence 78, Appl
39	17	0.8	159	4	US-08-639-075A-78	Sequence 78, Appl
40	17	0.8	159	4	US-09-012-411-78	Sequence 78, Appl
41	17	0.8	159	4	US-09-012-692-78	Sequence 78, Appl
42	17	0.8	159	4	US-08-906-613-78	Sequence 78, Appl
43	17	0.8	159	5	PCT-US95-14442A-78	Sequence 78, Appl
44	17	0.8	553	4	US-09-227-357-94	Sequence 94, Appl
45	17	0.8	558	2	US-08-896-385-5	Sequence 5, Appl

ALIGNMENTS

RESULT 1
US-08-936-165A-192/c
Sequence 192, Application US/08936165A
Patent No. 6348582
GENERAL INFORMATION:
APPLICANT: Black, Michael
APPLICANT: Barnham, Martin
APPLICANT: Hodgson, John
APPLICANT: Knowles, David
APPLICANT: Lonetto, Michael
APPLICANT: Nicholas, Richard
APPLICANT: Pratt, Julie
APPLICANT: Reichard, Richard
APPLICANT: Rosenberg, Martin
APPLICANT: Ward, Judith
TITLE OF INVENTION: No. 6348582a1 Prokaryotic Polynucleotides,
TITLE OF INVENTION: Polypeptides and Their Uses
NUMBER OF SEQUENCES: 534
CORRESPONDENCE ADDRESS:
ADDRESSEE: SmithKline Beecham Corporation
STREET: 709 Sweden Road
CITY: King of Prussia
STATE: PA
COUNTRY: USA
ZIP: 19406 0949
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FASTSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/936,165A
FILING DATE: 24-SEP-1997
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/027,032
FILING DATE: 24-SEP-1996
ATTORNEY/AGENT INFORMATION:
NAME: Gimm, Howard R
REGISTRATION NUMBER: 48,891
REFERENCE/DOCKET NUMBER: P50549
TELECOMMUNICATION INFORMATION:
TELEPHONE: 610-270-4478
FAX: 610-270-5090
FILES:
INFORMATION FOR SEQ ID NO: 192:
SEQUENCE CHARACTERISTICS:
LENGTH: 479 base pairs
TYPE: nucleic acid
STRANDNESS: single

GENERAL INFORMATION:
APPLICANT: Cohen, Daniel
APPLICANT: Blumenfeld, Maria
APPLICANT: Ilya, Chumakov
TITLE OF INVENTION: Prostate cancer gene
FILE REFERENCE: GENSET-018CPI
CURRENT APPLICATION NUMBER: US/09/218,207
CURRENT FILING DATE: 1998-12-22
EARLIER APPLICATION NUMBER: 08/996,306
EARLIER FILING DATE: 1997-12-22
EARLIER APPLICATION NUMBER: 60/099,658
EARLIER FILING DATE: 1998-09-09
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SOFTWARE: Patent.pm
SEQ ID NO: 183
LENGTH: 37950
TYPE: DNA
ORGANISM: Mus musculus
FEATURE:
NAME/KEY: exon
LOCATION: 5259..5328
OTHER INFORMATION: exon2
FEATURE:
NAME/KEY: exon
LOCATION: 12675..12791
OTHER INFORMATION: exon3
FEATURE:
NAME/KEY: exon
LOCATION: 14621..14710
OTHER INFORMATION: exon4
FEATURE:
NAME/KEY: exon
LOCATION: 19822..19912
OTHER INFORMATION: exon5
FEATURE:
NAME/KEY: exon
LOCATION: 21789..21950
OTHER INFORMATION: exon6
FEATURE:
NAME/KEY: exon
LOCATION: 23387..23510
OTHER INFORMATION: exon7
FEATURE:
NAME/KEY: exon
LOCATION: 25520..26016
OTHER INFORMATION: exon8
US-09-218-207-183

Query Match 0.98; Score 19; DB 4; Length 37950;
Best Local Similarity 100.0%; Pred. No. 7.9;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1320 cctcttattactgtaaaat 1338
Db 1077 cctcttattactgtaaaat 1095
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RESULT 5
US-08-744-419-3
Sequence 3, Application US/08744419
Patent No. 6274342
GENERAL INFORMATION:
APPLICANT: Guillerrez-Ramos et al.
TITLE OF INVENTION: Monocyte Chemotactic Protein 5 (MCP-5) Molecules
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: LAHIVE & COCKFIELD
STREET: 60 State Street, suite 510
CITY: Boston
STATE: Massachusetts
COUNTRY: USA

ZIP: 02109-1875
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
CREATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/744,419
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: U.S. Provisional
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Atwood, Beth E.
REGISTRATION NUMBER: 35,430
REFERENCE/DOCKET NUMBER: M10-008
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617)227-7400
TELEFAX: (617)227-5941
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 315 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-08-744-419-3

Query Match 0.98; Score 18; DB 4; Length 315;
Best Local Similarity 100.0%; Pred. No. 23;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1824 jctgaagaactacagacg 1841
Db 138 actgaagaactacagacg 155
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RESULT 6
US-08-744-419-1
Sequence 1, Application US/08744419
Patent No. 6274342
GENERAL INFORMATION:
APPLICANT: Guillerrez-Ramos et al.
TITLE OF INVENTION: Monocyte Chemotactic Protein 5 (MCP-5) Molecules
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: LAHIVE & COCKFIELD
STREET: 60 State Street, suite 510
CITY: Boston
STATE: Massachusetts
COUNTRY: USA
ZIP: 02109-1875
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/744,419
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: U.S. Provisional
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Atwood, Beth E.
REGISTRATION NUMBER: 35,430
REFERENCE/DOCKET NUMBER: M10-008
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617)227-7400
TELEFAX: (617)227-5941

FEATURE:
NAME/KEY:
LOCATION:
IDENTIFICATION METHOD: DNA sequencing and restriction analysis
OTHER INFORMATION: The encoded product of nucleotide SEQ ID NO: 1 is the human
PUBLICATION INFORMATION:
AUTHORS: R. Lomneda et al
TITLE: Cloning and sequencing of a cDNA encoding human
TITLE: milk beta-casein.
JOURNAL: Federation European Biochemical Society Letters
VOLUME: 269
ISSUE:
PAGES: 153 - 156
DATE: 1990
DOCUMENT NUMBER:
FILING DATE:
PUBLICATION DATE:
RELEVANT RESIDUES IN SEQ ID NO:
US-08-249-554-1

Query Match 0.9%; Score 18; DB 1; Length 1065;
Best Local Similarity 100.0%; Pred. No. 24;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 cagaagctgtgctgaa 19
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DB 192 CAGAGCTGTGCTGAA 175

RESULT 9

US-08-249-555A-1/c

Sequence 1, Application US/08249555A

Patent No. 5538952

GENERAL INFORMATION:

APPLICANT: Mokerji, P.

APPLICANT: Seo, A.

APPLICANT: Anderson, S.

APPLICANT: Schaller, J.

TITLE OF INVENTION: Inhibition of Infection of Mammalian Cells by Respiratory Sync

NUMBER OF SEQUENCES: 5

CORRESPONDENCE ADDRESS:

ADDRESSEE: Lonnie R. Drayer

ADDRESSEE: Ross Products Division

ADDRESSEE: Abbott Laboratories

STREET: 625 Cleveland Avenue

CITY: Columbus

STATE: Ohio

COUNTRY: United States

ZIP: 43215

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 inch, 1.44 Mb storage (B)COMPUTER: Apple Macintosh

OPERATING SYSTEM: Macintosh System 7.1(D)SOFTWARE: ClarisWorks 1.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/249,555A

FILING DATE: 26-MAY-1994

CLASSIFICATION: 514

PRIOR APPLICATION DATA: No. 5538952 applicable

TELECOMMUNICATION INFORMATION:

TELEPHONE: (614) 624-3774

TELEFAX: (614) 624-3074

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1065 base pairs

TYPE: Nucleic acid

STRANDEDNESS: Single

TOPOLOGY: Unknown

MOLECULE TYPE: Cloned cDNA representing the product of a human

DESCRIPTION: Human milk aeta-casein

HYPOTHEICAL:

ANTI-SENSE:

FRAGMENT TYPE:
ORIGINAL SOURCE: Human
ORGANISM: Homo sapiens
STRAIN:
INDIVIDUAL ISOLATE:
DEVELOPMENTAL STAGE: Adult
LAPOTYPE:
TISSUE TYPE: Mammary gland
CELL TYPE:
CELL LINE:
ORGANELLE:
IMMEDIATE SOURCE: Human Mammary Gland
LIBRARY:
CLONE:
POSITION IN GENOME:
CHROMOSOME/SEGMENT:
MAP POSITION:
UNITS:
FEATURE:
NAME/KEY:
LOCATION:
IDENTIFICATION METHOD: DNA sequencing and restriction analysis
OTHER INFORMATION: The encoded product of nucleotide SEQ ID NO: 1 is the human
PUBLICATION INFORMATION:
AUTHORS: R. Lomneda et al
TITLE: Cloning and sequencing of a cDNA encoding human
TITLE: milk beta-casein.
JOURNAL: Federation European Biochemical Society Letters
VOLUME: 269
ISSUE:
PAGES: 153 - 156
DATE: 1990
DOCUMENT NUMBER:
FILING DATE:
PUBLICATION DATE:
RELEVANT RESIDUES IN SEQ ID NO:
US-08-249-555A-1

Query Match 0.9%; Score 18; DB 1; Length 1065;
Best Local Similarity 100.0%; Pred. No. 24;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 cagaagctgtgctgaa 19
|||||
DB 192 CAGAGCTGTGCTGAA 175

RESULT 10

US-08-249-584-1/c

Sequence 1, Application US/08249584

Patent No. 5643880

GENERAL INFORMATION:

APPLICANT: Mokerji, P.

APPLICANT: Seo, A.

APPLICANT: Anderson, S.

APPLICANT: Harvey, L.

TITLE OF INVENTION: Product for Inhibition of Attachment of

NUMBER OF SEQUENCES: 5

CORRESPONDENCE ADDRESS:

ADDRESSEE: Lonnie R. Drayer

ADDRESSEE: Ross Products Division

ADDRESSEE: Abbott Laboratories

STREET: 625 Cleveland Avenue

CITY: Columbus

STATE: Ohio

COUNTRY: United States

ZIP: 43215

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 inch, 1.44 Mb storage

COMPUTER: Apple Macintosh

OPERATING SYSTEM: Macintosh System 7.1

```

SOFTWARE: Flatworks 1.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/249,584
FILING DATE: 26-MAY-1994
CLASSIFICATION: 424
PRIOR APPLICATION DATA: No. 564 0880 applicable
TRADITIONAL INVENTION INFORMATION:
INVENTOR: (614) 624-4774
TELEPHONE: (614) 624-4074
FAX: No. 564 0880
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1065 base pairs
TYPE: Nucleotide acid
STRANDNESS: Single
Topology: Unknown
MOLECULE TYPE: cloned cDNA representing the product of a human
MOLECULE TYPE: genomic DNA segment
DESCRIPTION: Human milk 'cald-casoin
HYDROTIC:
ANALYSIS:
FRAGMENT TYPE:
ORIGINAL SOURCE: Human
ORGANISM: Homo sapiens
STRAIN:
INDIVIDUAL ISOLATE:
DEVELOPMENTAL STAGE: Adult
HAPLOTYPE:
TISSUE TYPE: Mammary gland
CELL TYPE:
CELL LINE:
ORGANELLER:
IMMEDIATE SOURCE: Human Mammary gland
LIBRARY:
CLONING IN GENOME:
CHROMOSOME/SEQUENCE:
MAP POSITION:
UNIT:
FEATURE:
NAME/KEY:
IDENTIFICATION METHOD: DNA sequencing and restriction
IDENTIFICATION METHOD: analysis
OTHER INFORMATION: The encoded product of nucleotide
OTHER INFORMATION: SEQ ID NO:1 is the human milk protein.
OTHER INFORMATION:
PUBLICATION INFORMATION:
AUTHOR: R. Jomard et al
TITLE: Cloning and sequencing of a cDNA encoding human
TITLE: milk beta casein.
JOURNAL: Federation European Biochemical Society Letters
VOLUME: 259
ISSUE:
PAGES: 153-156
DATE: 1990
DOCUMENT NUMBER:
FILE NAME:
PUBLICATION DATE:
RELEVANT RESIDUES IN SEQ ID NO:
US 08 249 584-1

```

```

Query Match: 0.98; Score 18; LR 1; Length 1065;
Best Local Similarity: 100.0%; Prod. No. 24;
Method: 18; Conserved: 0; Mismatches: 0; Indels: 0; Gaps: 0;
97 2 complementarity 19
100 100% CAGAGAGTCTGAGGAA 175
RESULT 11

```

```

US 08 734-792-1/6
Sequence 1, Application US/08/44792
Patent No. 5707968
GENERAL INFORMATION:
APPLICANT: MOKER, J. P.
APPLICANT: Soc. A.
APPLICANT: Anderson, S.
APPLICANT: Harvey, L.
TITLE OF INVENTION: Inhibition of Attachment of H. Influenzae
TITLE OF INVENTION: to Human Cells
NUMBER OF SEQUENCES: 5
CROSSREFERENCE ADDRESS:
ADDRESS: Lantate R. Dreyer
ADDRESS: Ross Products Division
ADDRESS: Abbott Laboratories
STREET: 625 Cleveland Avenue
CITY: Columbus
STATE: Ohio
COUNTRY: United States
ZIP: 43215
COMPUTER READABLE FORM:
SEQUIM TYPE: 4.5 inch, 1.44 Mb storage
COMPIER: Apple Macintosh
OPERATING SYSTEM: Macintosh System 7.1
SOFTWARE: Flatworks 1.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/734,792
FILING DATE: 26-MAY-1994
CLASSIFICATION: 514
PUBLICATION INFORMATION:
APPLICATION NUMBER: 08/249,556
FILING DATE: 26-MAY-1994
TELEPHONE: (614) 624-4774
FAX: (614) 624-4074
TELEPHONE: (614) 624-4074
FAX: No. 5707968
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1065 base pairs
TYPE: Nucleotide acid
STRANDNESS: Single
Topology: Unknown
MOLECULE TYPE: cloned cDNA representing the product of a human
MOLECULE TYPE: genomic DNA segment
DESCRIPTION: Human milk 'cald-casoin
HYDROTIC:
ANALYSIS:
FRAGMENT TYPE:
ORIGINAL SOURCE: Human
ORGANISM: Homo sapiens
STRAIN:
INDIVIDUAL ISOLATE:
DEVELOPMENTAL STAGE: Adult
HAPLOTYPE:
TISSUE TYPE: Mammary gland
CELL TYPE:
CELL LINE:
ORGANELLER:
IMMEDIATE SOURCE: Human Mammary gland
LIBRARY:
CLONING IN GENOME:
CHROMOSOME/SEQUENCE:
MAP POSITION:
UNIT:
FEATURE:
NAME/KEY:
IDENTIFICATION METHOD: DNA sequencing and restriction
IDENTIFICATION METHOD: analysis
OTHER INFORMATION: The encoded product of nucleotide
OTHER INFORMATION: SEQ ID NO:1 is the human milk protein, 'casein.
PUBLICATION INFORMATION:

```


AUTHORS: B. Lonnnerdal et al
TITLE: Cloning and sequencing of a cDNA encoding human
MILK beta-casein
JOURNAL: Federation European Biochemical Society Letters
VOLUME: 269
ISSUE:
PAGES: 153 - 156
DATE: 1990
DOCUMENT NUMBER:
PUBLICATION DATE:
RELEVANT RESIDUES IN SEQ ID NO:
US-08-734-792-1

Query Match 0.98; Score 18; DB 1; Length 1065;
Best Local Similarity 100.0%; Pred. No. 24;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 caagagctgtgctgaa 19
|||||
DB 192 CAGAGCTGTGCTGCAA 175

RESULT 12
US-08-078-090-1/c
Sequence 1, Application US/08078090
Patent No. 5739407
GENERAL INFORMATION:
APPLICANT: BERGSTROM, SVEN
APPLICANT: HERNELL, OLLE
APPLICANT: LOENNERDAL, BO
APPLICANT: HJALMARSSON, KARIN
APPLICANT: HANSSON, LENNART
APPLICANT: TOERNELL, JAN
APPLICANT: STROMQUIST, MATS
TITLE OF INVENTION: HUMAN BETA-CASEIN PROCESS FOR PRODUCING
TITLE OF INVENTION: IT AND USE THEREOF
NUMBER OF SEQUENCES: 24
CORRESPONDENCE ADDRESS:
ADDRESSEE: BROWDY AND NEIMARK
STREET: 419 SEVENTH STREET, N.W.
CITY: WASHINGTON
STATE: D.C.
COUNTRY: USA
ZIP: 20004
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/078-090
FILING DATE: 19930618
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/DK92/00236
FILING DATE: 19-AUG-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/DK91/00233
FILING DATE: 19-AUG-1991
ATTORNEY/AGENT INFORMATION:
NAME: COOPER, IVER P.
REGISTRATION NUMBER: 28,005
REFERENCE/DOCKET NUMBER: BERGSTROM2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202)628-5197
TELEFAX: (202)737-3528
TELEX: 248633
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1065 base pairs
TYPE: nucleic acid

STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: sig.peptide
LOCATION: 4..48
FEATURE:
NAME/KEY: CDS
LOCATION: 49..681
US-08-078-090-1

Query Match 0.98; Score 18; DB 1; Length 1065;
Best Local Similarity 100.0%; Pred. No. 24;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 caagagctgtgctgaa 19
|||||
DB 192 CAGAGCTGTGCTGCAA 175

RESULT 13
PCT-US95-03628-1/c
Sequence 1, Application PC/TUS9503628
GENERAL INFORMATION:
APPLICANT: Abbott Laboratories
TITLE OF INVENTION: Inhibition of Infection of Mammalian Cells by
TITLE OF INVENTION: Respiratory Syncytial Virus
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Lonnie R. Dryer
ADDRESSEE: ROSS Products Division
ADDRESSEE: Abbott Laboratories
STREET: 625 Cleveland Avenue
CITY: Columbus
STATE: Ohio
COUNTRY: United States of America
ZIP: 43215
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 4.5 inch, 1.44 MB storage
COMPUTER: IBM compatible
OPERATING SYSTEM: MS-DOS Version 6.21
SOFTWARE: WordPerfect Version 6.0a
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/03628
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/249,554
FILING DATE: 26-MAY 1994
APPLICATION NUMBER: US 08/249,555
FILING DATE: 26-MAY-1994
TELECOMMUNICATION INFORMATION:
TELEPHONE: (614) 624-3774
TELEFAX: (614) 624-3074
TELEX: None
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1065 base pairs
TYPE: Nucleic acid
STRANDEDNESS: Single
TOPOLOGY: Unknown
MOLECULE TYPE: Cloned cDNA representing the product of a
MOLECULE TYPE: human genomic DNA segment
DESCRIPTION: Human milk eta-casein
HYPOTHETICAL:
ANTI-SENSE:
FRAGMENT TYPE:
ORIGINAL SOURCE: human
ORGANISM: Homo sapiens
STRAIN:
INDIVIDUAL ISOLATE:
DEVELOPMENTAL STAGE: Adult

```

1 HAPLOTYPE:
2 TISSUE TYPE: Mammary gland
3 CELL TYPE:
4 CELL LINE:
5 ORGANISM:
6 IMMEDIATE SOURCE: Human Mammary Gland
7 LIBRARY:
8 CLONING:
9 POSITION IN GENOME:
10 CHROMOSOME/SEGMENT:
11 MAP POSITION:
12 UNITS:
13 FEATURE:
14 NAME/KEY:
15 LOCATION:
16 IDENTIFICATION METHOD: DNA sequencing and restriction analysis
17 OTHER INFORMATION: The encoded product of nucleotide SEQ ID
18 OTHER INFORMATION: No: 1 is the human milk protein, "casein."
19 PUBLICATION INFORMATION:
20 AUTHORS: H. Jomard, et al.
21 TITLE: Cloning and sequencing of a cDNA encoding human milk beta-
22 TITLE: casein.
23 JOURNAL: Federation European Biochemical Society Letters
24 VOLUME: 269
25 ISSUE:
26 PAGES: 154 - 156
27 DATE: 1990
28 LOCATION NUMBER:
29 FILING DATE:
30 PUBLICATION DATE:
31 RELEVANT RESIDUES IN SEQ ID NO:
32 PCT-US95 0628-1

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Query Match 0.98; Score 18; Ind 5; Length 1065;
Best Local Similarity 100.0%; Prod. No. 24;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
DB 192 CACACAGCTGGCTGGAA 175

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RESULT 14
PCT-US95 01789-1/c
Sequence 1; Application 19/01US9503789
GENERAL INFORMATION:
1 APPLICANT: Abbott Laboratories
2 TITLE OF INVENTION: Inhibition of Attachment of H. Influenzae to Human
3 TITLE OF INVENTION: Cells
4 NUMBER OF SEQUENCES: 5
5 CORRESPONDENCE ADDRESS:
6 ADDRESSEE: Louie R. Dwyer
7 ADDRESSEE: Ross Products Division
8 STREET: 620 Cleveland Avenue
9 CITY: Columbus
10 STATE: Ohio
11 COUNTRY: United States of America
12 ZITE: 44215
13 COMPUTER READABLE FORM:
14 MEDIA TYPE: Diskette, 4.5 inch, 1.44 MB storage
15 COMPUTER: IBM compatible
16 OPERATING SYSTEM: MS DOS Version 6.04
17 SOFTWARE: WordPerfect Version 6.04
18 CURRENT APPLICATION DATA:
19 APPLICATION NUMBER: 19/01US95/03789
20 FILING DATE:
21 CLASSIFICATION:
22 PRIOR APPLICATION DATA:
23 APPLICATION NUMBER: US 08/249,556
24 FILING DATE: 26 MAY 1994
25 APPLICATION NUMBER: US 08/249,584

```

```

1 FILING DATE: 26 MAY 1994
2 TELECOMMUNICATION INFORMATION:
3 TELEPHONE: (614) 624-4774
4 TELEFAX: (614) 624-4074
5 TELETYPE: None
6 INFORMATION FOR SEQ ID NO: 1:
7 SEQUENCE CHARACTERISTICS:
8 LENGTH: 1065 base pairs
9 TYPE: Nucleic acid
10 STRANDEDNESS: Single
11 TOPOL/KEY: Unknown
12 MOLECULE TYPE: cloned cDNA representing the product of a
13 MOLECULE TYPE: human genomic DNA segment
14 DESCRIPTION: Human milk eta-casein
15 HYDROLYTIC:
16 ANTI SENSE:
17 FRAGMENT TYPE:
18 ORIGINAL SOURCE: Human
19 ORGANISM: Homo sapiens
20 STRAIN:
21 INDIVIDUAL ISOLATE:
22 DEVELOPMENTAL STAGE: Adult
23 HAPLOTYPE:
24 TISSUE TYPE: Mammary gland
25 CELL TYPE:
26 CELL LINE:
27 ORGANISM:
28 IMMEDIATE SOURCE: Human Mammary Gland
29 LIBRARY:
30 CLONING:
31 POSITION IN GENOME:
32 CHROMOSOME/SEGMENT:
33 MAP POSITION:
34 UNITS:
35 FEATURE:
36 NAME/KEY:
37 LOCATION:
38 IDENTIFICATION METHOD: DNA sequencing and restriction analysis
39 OTHER INFORMATION: The encoded product of nucleotide SEQ ID
40 OTHER INFORMATION: No: 1 is the human milk protein, "casein."
41 PUBLICATION INFORMATION:
42 AUTHORS: H. Jomard, et al.
43 TITLE: Cloning and sequencing of a cDNA encoding human milk beta-
44 TITLE: casein.
45 JOURNAL: Federation European Biochemical Society Letters
46 VOLUME: 269
47 ISSUE:
48 PAGES: 154 - 156
49 DATE: 1990
50 LOCATION NUMBER:
51 FILING DATE:
52 PUBLICATION DATE:
53 RELEVANT RESIDUES IN SEQ ID NO:
54 PCT-US95 01789-1

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Query Match 0.98; Score 18; Ind 5; Length 1065;
Best Local Similarity 100.0%; Prod. No. 24;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
DB 192 CACACAGCTGGCTGGAA 175

```

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RESULT 15
US 08 546 024 5/c
Sequence 5; Application US/08596024
Patent N. 5716847
GENERAL INFORMATION:
1 APPLICANT: Batty, Gerard F.
2 APPLICANT: Dwyer, Jan W.
3 APPLICANT: Kishoto, Ganesh M.

```

APPLICANT: Melton, Marcia L.
TITLE OF INVENTION: Expression of Sucrose Phosphorylase in
TITLE OF INVENTION: Plants
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Charles E. Cohen, Monsanto Company, R44F
STREET: 700 Chesterfield Parkway No. 5716837th
CITY: St. Louis
STATE: Missouri
COUNTRY: USA
ZIP: 63198
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn release #1.0, Version #1.30
CURRENT APPLICATION DATA: US/08/596.024
APPLICATION NUMBER: US/08/596.024
FILING DATE:
CLASSIFICATION: 415
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/386,860
FILING DATE: 10-FEB-1995
ATTORNEY/AGENT INFORMATION:
NAME: Cohen, Charles E.
REGISTRATION NUMBER: 34,565
REFERENCE/DOCKET NUMBER: 38-21(13567)A
TELECOMMUNICATION INFORMATION:
TELEPHONE: (314)537-6224
TELEFAX: (314)537-6047
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 1446 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-596-024-5

Query Match 0.98; Score 18; DB 1; Length 1446;
Best Local Similarity 100.0%; Pred. No. 24;
Matches 18; Conservative 0; Mismatches 0; Indels 0;
Gaps 0;
QY 459 gctttcttgatcct 476
TTTTTTTTTTTTTT
DB 304 GCTTTCTTGATCCT 287

Search completed: July 30, 2002, 05:14:10
Job time: 4662 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 30, 2002, 03:59:49 ; Search time 243.92 Seconds

(without alignments)
14556.326 Million cell updates/sec

Title: US-09-932-678-1

Perfect score: 2068
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Scoring table:
Gapop 60.0 , Gapext 60.0

Searched: 1736436 seqs, 858457221 residues

Word size : 0

Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : N.Geneseq_032802.*

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24: /SIDSL/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1869	90.4	2040	21	Human RNA polymerase I
2	920	44.5	1418	21	Human ORF2492
3	842	40.7	1498	22	Human full-length
4	691	33.4	1423	20	Human secretory mo
5	292	14.1	341	22	EST clone D1349.
6	269	13.0	3169	23	DNA encoding novel
7	232	11.2	1461	22	Human transcriptio
8	230	11.1	2103	21	Human cancer assoc
9	229	11.1	2991	23	DNA encoding novel

10	229	11.1	2991	23	AA591992	DNA encoding novel
11	223	10.8	776	23	AA592253	DNA encoding novel
12	215	10.4	3175	23	AA592254	DNA encoding novel
13	208	10.1	34269	22	AAK68677	Human immune/hama
14	208	10.1	34269	22	AAK85168	Human immune/hama
15	198	9.6	2410	23	AA592255	DNA encoding novel
16	196	9.5	437	20	AAK51663	Human secreted pro
17	190	9.2	396	23	AA580954	DNA encoding novel
18	190	9.2	396	23	AA580954	DNA encoding novel
19	190	9.2	396	23	AA580954	DNA encoding novel
20	190	9.2	396	23	AA580954	DNA encoding novel
21	187	9.0	17803	22	AAK68676	Human immune/hama
22	182	8.8	498	23	AA592247	DNA encoding novel
23	180	8.7	349	21	AAK77533	Human ORF2492
24	176	8.5	9259	22	AAK68674	Human immune/hama
25	176	8.5	23815	22	AAK68678	Human immune/hama
26	176	8.5	23815	22	AAK68678	Human immune/hama
27	175	8.5	585	23	AA592248	DNA encoding novel
28	174	8.4	1017	23	AA587646	DNA encoding novel
29	139	6.7	655	23	AA580955	DNA encoding novel
30	139	6.7	1260	23	AA587643	DNA encoding novel
31	133	6.4	627	23	AA587640	DNA encoding novel
32	117	5.7	213	21	AAK19091	Human secreted pro
33	110	5.3	3410	22	AAK85167	Human immune/hama
34	99	4.8	560	23	AA591991	DNA encoding novel
35	91	4.4	1840	22	AAK94394	Human full-length
36	80	3.9	405	22	AAK56711	Human immune/hama
37	76	3.7	414	22	AAK6215	Human breast cell
38	76	3.7	414	22	AAK6215	Human breast cell
39	76	3.7	414	22	AAK6215	Human breast cell
40	76	3.7	414	22	AAK6215	Human breast cell
41	76	3.7	414	22	AAK6215	Human breast cell
42	76	3.7	414	22	AAK6215	Human breast cell
43	76	3.7	414	22	AAK6215	Human breast cell
44	76	3.7	414	22	AAK6215	Human breast cell
45	75	3.6	95	22	ABA51325	Human breast cell

ALIGNMENTS

RESULT 1
ID AA598384 standard; cDNA: 2040 bp.
AC AA598384;
XX
XX 08-FEB-2001 (first entry)
XX
XX Human RNA polymerase I transcription factor TIF-1A cDNA.
DE
XX RNE polymerase I transcription factor TIF-1A; antitumor; treatment;
KW antiproliferative; cell proliferation; cancer; tissue regeneration; S.
XX
XX Hcno sapiens.
OS
XX
XX W030055316-A1.
XX
XX 21-SEP-2000.
XX
XX 08-MAR-2000; 2000WO-DE00767.
XX
XX 17-MAR-1999; 99DE-1011992.
XX
XX (DEKRR-) DEUT KRENSFORCHUNGSZENTRUM.
XX
XX Gramin I, Vintron M;
XX
XX WP1: 2000-587527/55.
XX
XX P-1SDS; AAB10936.
XX
XX New DNA encoding the transcription factor TIF-1A, useful for prevention
PT of treating diseases associated with abnormal cell proliferation.

QY	1607	tttggcaatccaaataatgacagctctgtcttcgtcaacccatcattgaaqaaanaa	1666
Dp	1578	ttgttgaacaccaaataatgacacgtctgtcttcgtcaacccatcattgaaqaaanaa	1637
QY	1667	tcgtccaaatgtgtgcacatgcatlaaqaatgacccgttgaqaqaacttcagtcgaatctgac	1726
Dp	1638	tcgtccaaatgtgtgcacatgcatlaaqaatgacccgttgaqaqaacttcagtcgaatctgac	1697
QY	1727	aaacccgtctgaaacacttcttccccccttgaacccctgatatgtctgaaagacccaagaatc	1786
Dp	1698	aaacccgtctgaaacacttcttccccccttgaacccctgatatgtctgaaagacccaagaatc	1757
QY	1787	cattgacacctattatcaggtatggtgaagaaatgattgtctgaagaaagctacagtgattcda	1846
Dp	1758	cattgacacctattatcaggtatggtgaagaaatgattgtctgaagaaagctacagtgattcda	1817
QY	1847	gaaacccatgaaadaaagacatattggaagacgaaatatgatacttctgaaagacgaat	1906
Dp	1818	gaaacccatgaaadaaagacatattggaagacgaaatatgatacttctgaaagacgaat	1877
QY	1907	gcccagaatcatatgcgtgatttggaatcaaacccaaccccttggacacgattttccgaag	1966
Dp	1878	gcccagaatcatatgcgtgatttggaatcaaacccaaccccttggacacgattttccgaag	1937
QY	1967	tccttcaagctagtgaggctccccaacccgttgtgtgacatgcaaacccagtcctccttaagc	2026
Dp	1938	tccttcaagctagtgaggctccccaacccgttgtgtgacatgcaaacccagtcctccttaagc	1997
QY	2027	gcacaaattttgagctgaagatgtaacatttggaattcccat	2086
Dp	1998	gcacaaattttgagctgaagatgtaacattttggaattcccat	2039

RESULT 2

ID AAC76937 standard; cDNA; 1418 BP.

AC AAC76937;

DT 08-FEB-2001 (first entry)

Human ORF2492 polynucleotide sequence SEQ ID NO:4983.

KW Human: open reading frame; ORF; detection; cytosolic; hepatotropic;
 KW vulnare; antiparasitic; antiparkinsonian; neurotropic; neuroprotective;
 KW anticonvulsant; osteopathic; antiarrhythmic; immunosuppressant; cardiac
 KW immunostimulant; thrombolytic; coagulant; vasotropic; antiatheric;
 KW hypotensive; dermatological; immunosuppressive; antiinflammatory;
 KW antiviral; antibacterial; antifungal; antirheumatic; antihypert;
 KW antiandrogen; gene therapy; cancer; proliferative disorder; hypertension;
 KW neurodegenerative disorder; osteoarthritis; graft vs host disease;
 KW cardiovascular disease; diabetes mellitus; hypothyroidism; SCID; AIDS;
 KW cholesterol ester storage; systemic lupus erythematosus; infection;
 KW severe combined immunodeficiency; malaria; autoimmune disorder; asthma;
 KW allergy; aplastic anaemia; nocturnal haemoglobinuria; burn; wound;
 KW bone disease; catallase damage; antiinflammatory disease; coagulation;
 KW thrombosis; contraceptive; ss.

US Homo sapiens.

PN W0200058473-A2.

PD 05-OCT-2000.

PF 31-MAR-2000; 2000WC-US08621.

PR 31-MAR-1999; 99US-0127607.

PR 05-APR-1999; 9905-0127728.

XX

XX

PI Shimkets RA, Leach M;

DR WPI; 2000-602362/57.

XX

PT useful for treating e.g. cancers, proliferative disorders,

XXXX

XX

CC AAC77444 to AAC77506 encode the proteins given in ABR0247 to ABR43894
CC which represent the human ORFX open reading frames 1 to 361. The ORFX
CC sequences have activities such as: cytostatic; hepatotropic; vulvotropic;
CC antiproliferative; antiparkinsonian; neurotropic; neuroprotective;
CC osteoplastic; anticonvulsant; antitachycardic; immunosuppressant;
CC immunostimulant; cardiant; thrombolytic; coagulant; vasoregret;
CC antidiabetic; hypotensive; dermatolytic; immunosuppressive;
CC antitumor; antiproliferative; antiviral; antifungal; antithrombotic;
CC antihypertensive; and antineoplastic. The sequences can be used for determining
CC the presence of or predisposition to, or prevention or treating
CC pathological conditions associated with an ORFX associated disorder. The
CC nucleic acids can be used to express ORFX proteins in gene therapy
CC vectors. The proteins and nucleic acids may be used to treat cancers,
CC proliferative disorders, neurodegenerative disorders, osteoarthritis,
CC graft vs host disease, cardiovascular disease, diabetes mellitus,
CC hypertension, hypothyroidism, cholesterol ester storage, systemic lupus
CC erythematosus, severe combined immunodeficiency (SCID), AIDS, viral,
CC bacterial or fungal infection, malaria, autoimmune disorders, asthma,
CC allergies, aplastic anaemia, burns, wounds, bone and cartilage damage,
CC neurological haemorrhagia, antinflammatory disease, to enhance
CC coagulation; to inhibit thrombosis; and as a contraceptive.

Sequence 1418 bp; 422 A; 275 C; 318 G; 403 T; 0 other.

Query March 44.58; Score 920; DB 21; Length 1418.

```
Matches 1120; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
```

Db 113 ccttaggcataaatctctggagcttatcttgaagaacttaactcaacttggatgttgaatc 172

QY 841 catccgcacqqlat.lgaagqlccl.gaaagaacacq'uar'laaactlqlqlqqarraq 501

501 attcaccgaagatttgllaatatqqatgaagclgaagaaactqadccatgaatcaagc 560
561

Db 2.3 attccacgaaagccttcaatattgcatggaagacccaagaaacccaacatggaacaaagg 292

961 ctgtctcctgaacccgtctcgaaccagatctgtgtgcatctctatgaatccctga 102

Db 293 ctggtcctgaacggtcgcaccagatggtgcacccctgtagccgagcgcctggaacatcctga 352

QY 1021 tctctttgatttgcctacatgaagatctctctatgtaagatggtaaggttgaataacg 108

Db 353 tgccttgccttgcctacatgaaqatgctcctatgtaaatgaaatcga 412

Qy 1061 accaaacaaqatctatatccqqaacqatbaaacatcttgaacaaactccqtqqa 114

Db 433 gcaaaccaaaqat.ctat.t.cq'qa'ctqal.aa'at.ct.t.qacaaat.ct.t.q.t.q'cc'a 472

Qy 1141 accatqctccatqlacat.t.t.t.c.alq.t.t.t.c.c.c.c.l.al.aq.t.t.c.a.a.t.t.q.q.q.t 120

Db 473 ccacatgcctccatgtaacatttllcagtttaacccctgtaatttccaaattcccat 512

QY 1201 ccqcaqacatlllllqgaacatctclqgaadaallqcaqacccaqladlcttqcca 120

5:3 tgcgaagcattttggaacatcttggaaaattg'aagaccc'aatlaattcttgc'a 592

Qy 1261 tcatcagwagqetqclqaaat tatat t ggaagctttt t qcaaqac t aaat t at t c 132

|||||
Db AACATCTTGCAACATCTCTTGTCCCAACATGCTCTCTGCAATATCAATTTTATG 440
QY 1173 TTTAATCTGTAGTTTCAATATGATTCGCAAGGACATTTTGAACATCTGAA 1212
Db 441 TTAACTCTGATTTCAATATGATTCGCAAGGACATTTTGAACATCTGAA 500
QY 1233 AATATGCAAGGACATTTCAATATGATTCGCAAGGACATTTTGAACATCTG 1292
Db 501 AATATGCAAGGACATTTCAATATGATTCGCAAGGACATTTTGAACATCTG 560
QY 1293 AGCTTTTGCAAGGACATTTCAATATGATTCGCAAGGACATTTTGAACATCT 1352
Db 561 AGCTTTTGCAAGGACATTTCAATATGATTCGCAAGGACATTTTGAACATCT 620
QY 1353 TTGATTAATCTGTAGTTTCAATATGATTCGCAAGGACATTTTGAACATCT 1412
Db 621 TTGATTAATCTGTAGTTTCAATATGATTCGCAAGGACATTTTGAACATCT 680
QY 1413 GATGTTCT 1472
Db 681 GATGTTCT 740
QY 1473 TTAAACCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGAC 1532
Db 741 TTAAACCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGAC 800
QY 1533 CTGAATTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGG 1592
Db 801 CTGAATTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGG 860
QY 1593 GTGATTAATCTGTAGTTTCAATATGATTCGCAAGGACATTTTGAAGGAC 1652
Db 861 GTGATTAATCTGTAGTTTCAATATGATTCGCAAGGACATTTTGAAGGAC 920
QY 1653 AT 1654
Db 921 AT 922

RESULT 4
AAS01562
ID AAS01562 standard; cDNA: 1423 bp.
XX
AC AAS01562;
XX
DT 18-JUL-2001 (first entry)
XX
DE Human secretory molecule cDNA sp1m #52.
XX
KW Human; secretory molecule; sp1m; SPTM; library screening; gene therapy;
KW cell signaling; cell proliferative disorder; atherosclerosis; cancer;
KW immune system disorder; AIDS; neurological disorder; Alzheimer's disease;
KW nervous system disorder; mental retardation; developmental disorder;
KW neuromuscular disorder; microarray; Incyte ID number 4287452dec; ss.
XX
OS Homo sapiens.
XX
PN W0200123558-A2.
XX
PD 05-APR-2001.
XX
PF 19-SEP-2000; 2000WO-US25610.
XX
PR 28-SEP-1999; 99US-0156624.
PR 28-SEP-1999; 99US-0156625.
PR 02-DEC-1999; 99US-0168611.
PR 02-DEC-1999; 99US-0168613.
PR 02-DEC-1999; 99US-0168614.
XX
PA (INCYTE) INCYTE GENOMICS INC.
XX
PI Hodson DM, Lincoln SE, Kusso FD, Spiro PA, Banville SC;

PI Brucher SR, Dutoir GE, Cohen BJ, Rosen RH, Shah P, Chakrap MS;
PI Hillman JL, Jones AL, Yu JY, Greenwalt LB, Panzer SR;
PI Roseberry AM, Wright RJ, Chen W, Liu TF, Yap PE, Stockdreyer IK;
PI Amshay S, Fong WT;
DR WPI: 2001-258134/26.
XX
PF New secretory polynucleotides (sp1m) and the polypeptides they encode,
PF for use in inducing antibodies and screening libraries of compounds.
XX
PS Claim 1: Page 152: 161pp; English.
XX
CC The present sequence for human secretory molecule cDNA sp1m #52
CC (Incyte ID number 4287452dec) is 1 of 63 novel sp1m cDNA sequences
CC (AAS01511-AAS01573) which encode for the secretory polypeptides SPTM. The
CC sp1m polynucleotides are useful for screening a compound for
CC effectiveness in altering expression of a target polynucleotide, where
CC the target polynucleotide comprises sp1m. Sp1m is also useful in a method
CC for assessing the toxicity of a test compound. Sp1m and its fragments or
CC complementary sequences, may be used to identify the presence of and/or
CC determine the degree of similarity between two nucleic acid sequences.
CC Sp1m can also be used for a variety of diagnostic and therapeutic
CC purposes, e.g., diagnosing a particular condition, disease or disorder
CC associated with cell signaling, such conditions include cell
CC proliferative disorders such as atherosclerosis, and cancers including
CC leukemia, an immune system disorder e.g., acquired immunodeficiency
CC syndrome (AIDS), a neurological disorder such as epilepsy or Alzheimer's
CC disease, nutritional and metabolic disease of the nervous system, mental
CC retardation and other developmental disorders, and muscular dystrophy and
CC other neuromuscular disorders. Sp1m can also be used to design probes
CC useful in diagnostic assays, which may be used to monitor the progress of
CC conditions or disorders associated with abnormal levels of expression of
CC sp1m. In addition sp1m encoding SPTM may be used for somatic or germ-line
CC gene therapy, for inducing antibodies, or in microarrays.
XX
SO Sequence 1423 bp; 417 A; 247 C; 320 G; 439 T; 0 other;

Query Match 44.4% Score 691; DH 22; Length 1423;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 691; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 169 TTTTCAATCT 248
Db 244 TTTTCAATCT 284
QY 249 CTGAATTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGAC 308
Db 254 CTGAATTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGAC 344
QY 309 CTGAATTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGAC 368
Db 344 CTGAATTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGAC 404
QY 369 TACTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGACAT 428
Db 404 TACTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGACAT 464
QY 429 AGAATTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGACAT 488
Db 454 AGAATTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGACAT 524
QY 459 ACTGTTCT 548
Db 524 ACTGTTCT 584
QY 549 ATCATTAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGACAT 608
Db 584 ATCATTAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGACAT 644
QY 609 GCAATTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGACAT 668
Db 644 GCAATTTGCAAGGACATTTTGAAGGACATTTTGAAGGACATTTTGAAGGACAT 704

PT	Novel transcription factor polypeptides, used to treat diseases
PI	associated with altered activity and expression of TRFX, and to screen
PI	for agents capable of modulating its activity -
XX	
PS	Claim 11: Page 299: 327bp: English.
XX	
CC	The present sequence is the coding sequence for a human transcription
CC	factor. The transcription factor and its coding sequence are useful in
CC	the diagnosis, treatment and prevention of diseases associated with
CC	altered expression of the transcription factor e.g. cell proliferative,
CC	autoimmune/inflammatory, neurological and developmental disorders. A
CC	number of specific disorders/diseases are given in the specification,
CC	including: arteriosclerosis, cirrhosis, hepatitis, cancers, AIDS,
CC	allergies, anaemia, asthma, autoimmune thyroiditis, bronchitis, atopic
CC	dermatitis, diabetes mellitus, emphysema, Goodpasture's syndrome, gout,
CC	grave's disease, multiple sclerosis, osteoarthritis, pancreatitis,
CC	psoriasis, rheumatoid arthritis, systemic lupus erythematosus, ulcerative
CC	colitis, uveitis, Alzheimer's disease, Huntington's disease, Parkinson's
CC	disease, stroke, and viral, bacterial, fungal and protozoal infections.
XX	
SO	Sequence 1461 BP: 413 A: 335 G: 298 G: 415 T: 0 other:
QY	Query Match 11.2% Score 232: 108 22: Length 1461:
Db	Best Local Similarity 96.6%: Pred. No. 2.8e-101:
Matches: 282: Conservative 0: Mismatches 1: Indels 0: Gaps 0:	
QY	1339 catgctaaatcttgaatgaatgctgcacatactaaactaaacacgaatcaggaa 1398
Db	6681 catgcctgaatccttttgctaaactgcgcgcacataactaaactaaacgaatcaggaa 740
QY	1359 caaagaacatctgcagatctgctctccatgagaaatcttaactgaactgcacacatgag 1458
Db	741 caaagcattctgcagatgctgctctccatgagcatttactgcacctgcacacatgag 800
QY	1459 tctaaccttcttttgaacacacacacacatttgacgcgaacacttgaagaagaatggc 1518
Db	801 tctaaccttcttttgaacacacacacacatttgacgcgaacacttgaagaagaatggc 860
QY	1519 agtacctcagatcttgaatttgaacagatagatgacgcacgttaactcctctgaaga 1578
Db	861 agtacctcagatcttgaatttgaacagatagatgacgcacgttaactcctctgaaga 920
QY	1579 ttgcctacccctcaatgataactttttgcgcacacacaa 1621
Db	921 ttgcctacccctcaatgataactttttgcgcacacacaa 963
RESULT 8	
AC	AACT779c
XX	ACACT7798 standard: cDNA: 2103 BP.
XX	ACACT7798:
XX	
DT	08-FEB-2001 (first entry)
XX	
DE	Human cancer associated gene sequence SEQ ID NO:192.
XX	
KM	Human: cancer associated gene: cancer antigen: detection: cancer:
KM	diancusi: cytosolic: proliferative: vlnucre: immunomodulatory:
KM	antiadhesive: antiasthmatic: antirheumatic: antiarthritis: antiviral:
KM	antiinflammatory: antithyroid: antiallergic: antibacterial: cardiac:
KM	dermatological: neuroprotective: thrombolytic: coagulant: neurologic:
KM	vasotropic: antipsoriatic: antiangiogenic: gene therapy: inflammation:
KM	immune disorder: haematopoietic cell disorder: autoimmune disorder:
KM	allergic reaction: graft versus host disease: organ rejection:
KM	haemostatic: thrombolytic: cardiovascular disorder: infection:
XX	neurological disease: drug screening: ss.
OS	Human sapiens.
XX	
PN	W0200055350-A1.
XX	

UY 561 ggcagatagatgttccatctcgaigaagatgata 600
 |||||||
 Db 850 ggcagatagatgttccatctcgaigaagatgata 889

RESULT 10

AAS91992
 ID AAS91992 standard; cDNA: 2991 BP.

AC AAS91992:

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #27796.

XX Human: chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX Homo sapiens.

FN W0200175067-A2.

PD 11-OCT-2001.

PE 30-MAR-2001: 2001WO-US08631.

PR 31-MAR-2000: 2000US-0540217.

PR 23-AUG-2000: 2000US-0649167.

XX (HYSE-) HYSEQ INC.

PI Dimaue RT, Liu C, Tang YT:

DR WPI: 2001-639362/73.

DR P-PSDB: ABG27805.

PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.

PS Claim 1; SEQ ID No 27796; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS6197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at http://wipo.int/pub/published_pat_sequences.

SO Sequence 2991 BP: 655 A; 853 C; 882 G; 601 T; 0 other;

Query Match 11.1%; Score 229; DB 23; Length 2991;
 Best Local Similarity 99.6%; Pred. No. 7,6e-100;
 Matches 279; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 321 gatgaccagatcaccactgagctgagatctccgtcttctctcagatgacttaccaga 480
 |||||||
 Db 610 gatgaccagatcaccactgagctgagatctccgtcttctctcagatgacttaccaga 669
 QY 381 gacttgaacacttaccatattatattaaatgcttgaatgaaatcacaac 440
 |||||||
 Db 670 gacttgaacacttaccatattatattaaatgcttgaatgaaatcacaac 729
 QY 441 gtaatgaagaatatttgccttttcttgaatgaaatcacaacatgctttctc 500
 |||||||
 Db 730 gtaatgaagaatatttgccttttcttgaatgaaatcacaacatgctttctc 789
 QY 501 agaccatgctcagatgagcttccatattgtctctcccaatgacatgaagaa 560
 |||||||
 Db 790 agaccatgctcagatgagcttccatattgtctctcccaatgacatgaagaa 849
 QY 561 ggcagatagatgttccatctcgaigaagatgata 600
 |||||||
 Db 850 ggcagatagatgttccatctcgaigaagatgata 889

RESULT 11

AAS9220
 ID AAS92253 standard; cDNA: 776 BP.

AC AAS92253:

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #28057.

XX Human: chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX Homo sapiens.

FN W0200175067-A2.

PD 11-OCT-2001.

PE 30-MAR-2001: 2001WO-US08631.

PR 31-MAR-2000: 2000US-0540217.

PR 23-AUG-2000: 2000US-0649167.

XX (HYSE-) HYSEQ INC.

PI Dimaue RT, Liu C, Tang YT:

DR WPI: 2001-639362/73.

DR P-PSDB: ABG28066.

PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.

PS Claim 1; SEQ ID No 28057; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations

PR 08-NOV-2000; 2000US-0246475.
 PR 08-NOV-2000; 2000US-0246476.
 PR 08-NOV-2000; 2000US-0246477.
 PR 08-NOV-2000; 2000US-0246478.
 PR 08-NOV-2000; 2000US-0246523.
 PR 08-NOV-2000; 2000US-0246524.
 PR 08-NOV-2000; 2000US-0246525.
 PR 08-NOV-2000; 2000US-0246526.
 PR 08-NOV-2000; 2000US-0246527.
 PR 08-NOV-2000; 2000US-0246528.
 PR 08-NOV-2000; 2000US-0246532.
 PR 08-NOV-2000; 2000US-0246609.
 PR 08-NOV-2000; 2000US-0246610.
 PR 08-NOV-2000; 2000US-0246611.
 PR 08-NOV-2000; 2000US-0246613.
 PR 17-NOV-2000; 2000US-0249207.
 PR 17-NOV-2000; 2000US-0249208.
 PR 17-NOV-2000; 2000US-0249209.
 PR 17-NOV-2000; 2000US-0249210.
 PR 17-NOV-2000; 2000US-0249211.
 PR 17-NOV-2000; 2000US-0249212.
 PR 17-NOV-2000; 2000US-0249213.
 PR 17-NOV-2000; 2000US-0249214.
 PR 17-NOV-2000; 2000US-0249215.
 PR 17-NOV-2000; 2000US-0249216.
 PR 17-NOV-2000; 2000US-0249217.
 PR 17-NOV-2000; 2000US-0249218.
 PR 17-NOV-2000; 2000US-0249244.
 PR 17-NOV-2000; 2000US-0249245.
 PR 17-NOV-2000; 2000US-0249264.
 PR 17-NOV-2000; 2000US-0249265.
 PR 17-NOV-2000; 2000US-0249297.
 PR 17-NOV-2000; 2000US-0249299.
 PR 17-NOV-2000; 2000US-0249300.
 PR 01-DEC-2000; 2000US-0250160.
 PR 01-DEC-2000; 2000US-0250391.
 PR 05-DEC-2000; 2000US-0251030.
 PR 05-DEC-2000; 2000US-0251988.
 PR 05-DEC-2000; 2000US-0256719.
 PR 06-DEC-2000; 2000US-0251479.
 PR 08-DEC-2000; 2000US-0251856.
 PR 08-DEC-2000; 2000US-0251868.
 PR 08-DEC-2000; 2000US-0251869.
 PR 08-DEC-2000; 2000US-0251989.
 PR 08-DEC-2000; 2000US-0251990.
 PR 11-DEC-2000; 2000US-0254097.
 PR 05-JAN-2001; 2001US-0259678.
 XX
 PA (HUMAN-) HUMAN GENOME SCI INC.
 XX
 P1 Roscon CA, Barash SC, Ruben SM;
 XX
 DR WPI: 2001-483426/52.
 XX
 PT Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
 PT useful for preventing, diagnosing and/or treating cancers and
 PT metastasis.
 XX
 PS Disclosure: SEQ ID NO 39980; 3071bp + Sequence listing: English.
 XX

AAK64702 encode the human immune/hematopoietic antigen (1)
 amino acid sequences given in AAK62170 to AAK61921. (1) have cytosolic
 activity, and can be used in gene therapy and vaccine production. (1)
 proteins and polynucleotides may be used in the prevention, diagnosis and
 treatment of diseases associated with inappropriate (1) expression. For
 example, they may be used to treat disorders associated with decreased
 expression by rectifying mutations or deletions in a patient's genome
 that affect the activity of (1) by expressing inactive proteins or to
 supplement the patient's own production of (1). Additionally, (1)
 polynucleotides may be used to produce the secreted (1), by inserting
 the nucleic acids into a host cell and culturing the cell to express the
 protein. (1) proteins and polynucleotides may be used to prevent,
 diagnose and treat immune/hematopoietic-related diseases, especially

CC cancers and cancer metastases of haematopoietic-derived cells. AAK64704
 CC to AAK67694 represent human immune/hematopoietic antigen genomic
 CC sequences from the present invention. AAK54942 to AAK54950 and AAK82169
 CC represent sequences used in the exemplification of the present invention.
 XX
 SQ Sequence 34269 BP; 9052 A; 6778 C; 7638 G; 10801 T; 0 other;
 Query Match 10.1%; Score 208; DR 22; Length 44269;
 Best Local Similarity 100.0%; Prod. No. 9.3e-90;
 Matches 208; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1861 aqacataatgaaatataaataatagatcttctgaagaagcaaatgcccgaatata 1920
 DB ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 DB 32384 aqacataatgaaatataaataatagatcttctgaagaagcaaatgcccgaatata 32444
 QY 1921 cctgattggatcacacaaatctcttgaacgcatctcgaatcttcaatattg 1980
 DB ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 DB 42444 cctgattggatcacacaaatctcttgaacgcatctcgaatcttcaatattg 42504
 QY 1961 taagctcccccacccgtttgtatcatgaaacccatccctctgaacacaaattatga 2040
 DB ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 DB 32504 taagctcccccacccgtttgtatcatgaaacccatccctctgaacacaaattatga 32564
 QY 2041 ctgaatattgaatttgaatttcccac 2068
 DB ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 DB 32564 ctgaatattgaatttgaatttcccac 32591
 RESULT 15
 AAK592255
 ID AAK592255 standard; cDNA; 2410 BP.
 XX
 AC AAK592255;
 XX
 DT 13-FEB-2002 (first entry)
 XX
 DE DNA encoding novel human diagnostic protein #28059.
 XX
 KW Human; chromosome mapping; gene mapping; gene therapy; forensics;
 KW food supplement; medical imaging; diagnosis; genetic disorder; ss.
 XX
 OS Homo sapiens.
 XX
 TN W0200175067-A2.
 XX
 PP 11-QCI-2001.
 XX
 PP 30-MAR-2001; 2001WO-0508631.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 PR 23-APR-2000; 2000US-0649167.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 P1 Dimaac RT, Liu C, Tang YT;
 XX
 DR WPI: 2001-639362/73.
 DR P-ISOB; AMO28068.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX
 PS Claim 1: SEQ ID No 28059; 103bp; English.
 XX
 CC The invention relates to isolated polynucleotide (1) and
 CC polypeptide (1) sequences. (1) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (1). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (1) is useful in gene therapy techniques

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 30, 2002, 03:10:00 ; Search time 2610.22 Seconds
(without alignments)
16579.475 Million cell updates/sec

Title: US-09-932-678-1

Perfect score: 2068
Sequence: 1 acagaagctatgctgagaa.....taacattggaattcccat 2068

Scoring table:

Gapop 60.0 , Gapext 60.0

Searched: 1797656 seqs, 10463268293 residues

Word size: 0

Total number of hits satisfying chosen parameters: 3595412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database:

GenBank:
1: gb_ba:*
2: gb_bg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vi:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_om:*
21: em_or:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pl:*
26: em_ro:*
27: em_sts:*
28: em_un:*
29: em_vi:*
30: em_hlg_hum:*
31: em_hlg_inv:*
32: em_hlg_other:*
33: em_hlgc_inv:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
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1	2068	100.0	2068	9	AF227156	AF227156 Homo sapi
2	1920	92.8	3756	9	HSZA272050	AJ272050 Homo sapi
3	1869	90.4	2040	6	AX036047	AX036047 Sequence
4	308	14.9	2493	9	AK055742	AK055742 Homo sapi
5	283	13.7	1177	9	BC009198	BC009198 Homo sapi
6	232	11.2	1461	6	AX274907	AX274907 Sequence
7	232	11.2	2627	9	BC006441	BC006441 Homo sapi
8	208	10.1	129473	2	AC040158	AC040158 Homo sapi
9	208	10.1	143900	2	AC092137	AC092137 Homo sapi
10	187	9.0	175651	2	AC007615	AC007615 Homo sapi
11	187	9.0	191496	2	AC009130	AC009130 Homo sapi
12	187	9.0	208008	2	AC009093	AC009093 Homo sapi
13	176	8.5	173166	9	AC092375	AC092375 Homo sapi
14	176	8.5	179150	2	AC017077	AC017077 Homo sapi
15	176	8.5	202004	9	BCAF001549	AF001549 Human Chr
16	158	7.6	143900	2	AC092137	AC092137 Homo sapi
17	125	6.0	170611	9	AC106788	AC106788 Homo sapi
18	123	5.9	187643	2	AC092562	AC092562 Papio ham
19	119	5.8	191496	2	AC009130	AC009130 Homo sapi
20	115	5.6	824	9	HSZA343405	AJ343405 Homo sapi
21	109	5.3	247331	2	AC097268	AC097268 Pan trogl
22	97	4.7	293	9	HS151B38	Z56854 H. sapiens C
23	92	4.4	179150	2	AC017077	AC017077 Homo sapi
24	79	3.8	196413	2	AC097269	AC097269 Pan trogl
25	69	3.3	78347	2	AC022642	AC022642 Homo sapi
26	54	2.6	78347	2	AC022642	AC022642 Homo sapi
27	47	2.3	108040	2	AC068150	AC068150 Homo sapi
28	47	2.3	171940	2	AC092119	AC092119 Homo sapi
29	47	2.3	173166	9	AC092375	AC092375 Homo sapi
30	47	2.3	316296	2	AC092285	AC092285 Homo sapi
31	42	2.0	821	9	HSZA343406	AJ343406 Homo sapi
32	36	1.7	405	11	G27379	G27379 human STS S
33	32	1.5	286	9	HS151B38	Z56854 H. sapiens C
34	30	1.5	15873	2	AC012547	AC012547 Homo sapi
35	30	1.5	192815	9	AC013444	AC013444 Homo sapi
36	26	1.3	461	11	G38441	G38441 SHC 58117
37	26	1.3	181088	2	AC097733	AC097733 Rattus no
38	24	1.2	54814	2	AC107769	AC107769 Mus muscu
39	24	1.2	53814	2	AC107769	AC107769 Mus muscu
40	22	1.1	737	9	HSZA37662	AJ37662 Homo sapi
41	22	1.1	38147	8	SPAC6812	Z98541 S. pombe chr
42	22	1.1	76416	9	AC005370	AC005370 Homo sapi
43	22	1.1	108793	9	AL137839	AL137839 Human DNA
44	22	1.1	146437	9	AC010489	AC010489 Homo sapi
45	22	1.1	155022	2	AC108176	AC108176 Bos tauru

ALIGNMENTS

RESULT 1
LOCUS AF227156
DEFINITION Homo sapiens RRM3 mRNA, complete cds.
ACCESSION AF227156
VERSION AF227156.1 GI:7670099
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 2068)
Moorefield,B., Greene,E.A. and Reeder,R.H.
RNA polymerase I transcription factor Rrm3 is functionally
conserved between yeast and human
Proc. Natl. Acad. Sci. U.S.A. 97 (9), 4724-4729 (2000)
MEDLINE 20243763
REFERENCE 2 (bases 1 to 2068)
Moorefield,B., Greene,E.A. and Reeder,R.H.
Direct Submission
Submitted (20-JAN-2000) Basic Sciences, Fred Hutchinson Cancer
Research Center, 1100 Fairview Avenue N., Seattle, WA 98109, USA
FEATURES
Location/Qualifiers

Db 1681 CAGTCATTAGGAGTACCGCTGAGAGAGACTGACGTGACGATCTGCACAAACCCGCTGACAA 1740

QY 1741 cctctctcccttgaatccctgtgtcctgaagagatcaaaatcattgactcattt 1800

Db 1741 ccttctcccttgaatccctgtgtcctgaagagatcaaaatcattgactcattt 1800

QY 1801 atcaagataaggagacatagatgctgaagagatcagagatcagagagacatcagaa 1860

Db 1801 ATCAGATATGGAGAGACATAGTCTGAGAGAGCTACAGAGGTTCAAGAAACCATGAGAAA 1860

QY 1861 aagacatagtagaagatcaagatgaatccttctgaagagagagatgcccccaatgata 1920

Db 1861 AGCAGATAGTGAAGATGAGATGATGATCTTCTGAAAGGCAAGTCCGCCAATGATA 1920

QY 1921 cctgtagtagatcagacacaaactccttctgaagagatcagagatccttctgaagatg 1980

Db 1921 cctgtagtagatcagacacaaactccttctgaagagatcagagatccttctgaagatg 1980

QY 1981 tggagctcccccagctgtgtgtagatcagacacaaactccttctgaagagatcagagatg 2040

Db 1981 TGGGCTCCGCCAGCTGTGTGATGATGATGATGATGATGATGATGATGATGATGATGATG 2040

QY 2041 ctgagatgtagatcagacacaaactccttctgaagagatcagagatccttctgaagatg 2068

Db 2041 CTCAGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 2068

RESULT 2

HSAG272050 3756 bp mRNA linear PRI 07-SEP-2000

LOCUS

DEFINITION Homo sapiens mRNA for transcription initiation factor 1A protein (TIF-1A gene).

ACCESSION AJ272050

VERSION AJ272050.1 GI:10046713

KEYWORDS TIF-1A; transcription initiation factor.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 3756)

AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

1 Boden, J., Hoffmann-Rohrer, U., Ross, W., Delius, H., Vinograd, M. and Grummt, I.

TITLE Cloning and functional characterization of transcription initiation factor TIF-1A, a growth-dependent regulator of ribosomal RNA synthesis

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 3756)

AUTHORS Boden, J.J.

TITLE Direct Submission

JOURNAL Submitted (07-FEB-2000) Boden J.J., Molekulare Biologie der Zelle 11, Deutsches Krebsforschungszentrum, INF.280, 69120 Heidelberg, GERMANY

FEATURES

Source

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/db_xref="taxon:9606"

23..3756 /gene="TIF-1A"

23..1978 /gene="TIF-1A"

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/codon_start=1

/evidence="experimental

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/db_xref="GI:10046714"

/translation="MAAPLITHTPLNDPDAASSAVKKLIGASRTGISNMALINDPENS

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ILKEGVNDSDDDDONLPANFTCHRALQILARYVSTPFWPLMLLEKSPVRS

ERTLECYNVALRIISVFPFLRHEILIELIKLIDVNASRGLIDAEIETATCGG

TSTFGLFNNDDEEETHEETKAGPERLDQVHVARLDIILMSLIVSYKDCYVNGK

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Db 1 cccgtcccttgaatcagacacaaactccttctgaagagatcagagatccttctgaagatg 60

QY 167 tggagctcccccagctgtgtgtagatcagacacaaactccttctgaagagatcagagatg 166

Db 61 tggagctcccccagctgtgtgtagatcagacacaaactccttctgaagagatcagagatg 120

QY 167 tggagctcccccagctgtgtgtagatcagacacaaactccttctgaagagatcagagatg 226

Db 121 tggagctcccccagctgtgtgtagatcagacacaaactccttctgaagagatcagagatg 180

QY 227 tggagctcccccagctgtgtgtagatcagacacaaactccttctgaagagatcagagatg 286

Db 181 tggagctcccccagctgtgtgtagatcagacacaaactccttctgaagagatcagagatg 240

QY 241 gttgag 346

Db 241 gttgag 300

QY 347 gttgag 406

Db 401 gttgag 360

QY 407 gttgag 466

Db 461 gttgag 420

QY 467 gttgag 526

Db 471 gttgag 480

QY 527 gttgag 586

Db 541 gttgag 540

QY 547 gttgag 646

Db 601 gttgag 600

QY 601 gttgag 660

Db 661 gttgag 720

QY 707 gttgag 766

Db 767 gttgag 720

QY 767 gttgag 826

Db 771 gttgag 780

QY 827 gttgag 886

Db 841 gttgag 840

QY 887 gttgag 946


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Db 757 GCAATTTTGACAT 772
RESULT 6
AX274907 1461 bp DNA Linear PAT 29-OCT-2001
LOCUS Sequence 172 from Patent WO0172777.
DEFINITION AX274907
ACCESSION AX274907
VERSION AX274907.1 GI:16547539
KEYWORDS
SOURCE
ORGANISM human.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
1 (sites)
Hillman, J.L., Baughn, M.R., Yue, H., Lal, P., Lu, D.A., Patterson, C.,
Arizumi, Y., Bandman, O., Tang, Y.F., Mathur, P., Shah, P., Au-Yang, J.,
and Reddy, R.
TITLE
Transcription factors
JOURNAL
Patent: WO 0172777-A 172 04-OCT-2001;
Incyte Genomics, Inc. (US)
FEATURES
Source
Location/Qualifiers
1..1461
/organism="Homo sapiens"
/db_xref="taxon:9606"
/note="Incyte ID No: 2646274CBI"
BASE COUNT 413 a 335 c 298 g 415 t
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Best local similarity 99.6% Pred. No. 1.2e-110;
Matches 282; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1339 catgctagatcttcttgatgaatgagatgacatataccttaaccagatcagaa 1398
Db 681 GATGCTAGATCTTGGTAACTGGCTGACATATACCTTAATAGCAGATTCGGAA 740
QY 1399 caaagcattcagatgattgctccatgagaccatttactcaagctcgaagctgt 1458
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QY 1459 tctaaccttcttctttagacaagaagcttcttgaagaaacctgaagaagatllgc 1518
Db 801 TCTACACCTTTGTTTATAGACACAGACGCTTTTGAGCGGAACCTGAAGAAGTTTGC 860
QY 1519 agtattcagagctgaattttagagagatagatgaagcagctaaatccctgaaga 1578
Db 861 AGATCTCTAGAGCTGAATTTTGGAGGAGATAGTATGATGAGCCAGCTAATCCCTGAAGA 920
QY 1579 tttagctccctcagtgatgaacttttttgcgaatcagaa 1621
Db 921 TTTCCTGCCCCCTAGTGTAACTTTTTCCTGCAATCAAGAA 963
RESULT 7
BC006441 2627 bp mRNA linear PRI 12-JUL-2001
LOCUS Homo sapiens, Similar to RNA polymerase 1 transcription factor
DEFINITION RRS3, clone MGC:13169 IMAGE:3351791, mRNA, complete cds.
ACCESSION BC006441
VERSION BC006441.1 GI:13623642
KEYWORDS
SOURCE
ORGANISM human.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
1 (bases 1 to 2627)
Strausberg, R.
TITLE
Direct Submission
JOURNAL
Submitted (09-APR-2001) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2550,

```

```

REMARKS
COMMENT
USA
NIH-MGC Project URL: http://mgc.ncl.nih.gov
Contact: MGC help desk
Email: mgc-help@mail.nih.gov
Tissue procurement: ATCC
cDNA Library Preparation: Rabin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LIN)
DNA Sequencing by: Genome Sequence Centre,
BC Cancer Agency, Vancouver, BC, Canada
info@bcsc.bc.ca
Steven Jones, Jennifer Asano, Ian Bosdet, Yaron Butterfield,
Susanna Chan, Readman Chiu, Chris Fjell, Erin Garland, Ran Guin,
Leticia Hsiao, Martin Krzywinski, Reta Kusche, Oliver Levy, Soo
Sen Lee, Victor Ling, Carrie Mathewson, Candice McLeavy, Steven
Ness, Pawan Pandoh, Anna-Lisa Prabhu, Parvaneh Saedi, Jacquelynne
Sechin, Duane Smalhus, Michael Smith, Lorraine Spencer, Jeff Stott,
Michael Thorpe, Miranada Tsai, Natasja van den Bosch, Jill Vardy,
George Yang, Scott Zuyderduyn, Marco Marra.
Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LIN at: http://image.jnl.gov
Series: IRAT Plate: 18 Row: d Column: 24
This clone was selected for full length sequencing because it
passed the following selection criteria: Hexamer frequency >8%
analysis.
FEATURES
Source
Location/Qualifiers
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/clone_lib="NIH-MGC_16"
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/note="Vector: pGB7"
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/codon_start=1
/product="Similar to RNA polymerase 1 transcription factor
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/protein_id="AAH06441.1"
/db_xref="GI:13623643"
/translat_id="MGFAVELEHMKMLDPSNVA11ROMA:NY10SLARAKE1SL
ITVKPCDLVLMHLILYNDSGTAKAPDVALHGPYSACVAVYTFVHKKJLHSL
NKEGLQYFOSINFERIYMSULNPKJLPLSVNFAMITMKKTGYEMW"
BASE COUNT 837 a 492 c 525 g 774 t
ORIGIN
Query Match 11.2% Score 232; DB 9; Length 2627;
Best local similarity 99.6% Pred. No. 1.2e-110;
Matches 282; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1339 catgctagatcttcttgatgaatgagatgacatataccttaaccagatcagaa 1398
Db 1643 GATGCTAGATCTTGGTAACTGGCTGACATATACCTTAATAGCAGATTCGGAA 1902
QY 1399 caaagcattcagatgattgctccatgagaccatttactcaagctcgaagctgt 1458
Db 1603 CAAAGCATTCCTCGATCTCTCTCCATGACCATTTTACTGAGCTGCAAGCTGTGT 1962
QY 1459 tctaaccttcttctttagacaagaagcttcttgaagaaacctgaagaagatllgc 1518
Db 1963 TCTACACCTTTGTTTATAGACACAGACGCTTTTGAGCGGAACCTGAAGAAGTTTGC 2022
QY 1519 agtattcagagctgaattttagagagatagatgaagcagctaaatccctgaaga 1578
Db 2023 AGATCTCTAGAGCTGAATTTTGGAGGAGATAGTATGATGAGCCAGCTAATCCCTGAAGA 2082
QY 1579 tttagctccctcagtgatgaacttttttgcgaatcagaa 1621
Db 2093 TTTCCTGCCCCCTAGTGTAACTTTTTCCTGCAATCAAGAA 2125
RESULT 8

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* arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

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1 1033: contig of 1033 bp in length
* 1034 1133: gap of unknown length
* 1134 2547: contig of 1414 bp in length
* 2548 2648: gap of unknown length
* 2648 4205: contig of 1558 bp in length
* 4206 4305: gap of unknown length
* 4306 6564: contig of 2259 bp in length
* 6565 9065: gap of unknown length
* 9066 9165: gap of unknown length
* 9166 12059: contig of 2894 bp in length
* 12060 12159: gap of unknown length
* 12160 19295: contig of 7136 bp in length
* 19296 19395: gap of unknown length
* 19396 25668: contig of 6273 bp in length
* 25669 25768: gap of unknown length
* 25769 31013: contig of 5245 bp in length
* 31014 31113: gap of unknown length
* 31114 36555: contig of 5442 bp in length
* 36556 45338: gap of unknown length
* 45339 45438: contig of 8683 bp in length
* 45439 52918: gap of unknown length
* 52919 53018: contig of 7480 bp in length
* 53019 65720: gap of unknown length
* 65721 65820: contig of 12702 bp in length
* 65821 76109: gap of unknown length
* 76110 76209: contig of 10289 bp in length
* 76210 85439: gap of unknown length
* 85440 85539: contig of 9230 bp in length
* 85540 101620: gap of unknown length
* 101621 101720: contig of 16081 bp in length
* 101721 122163: gap of unknown length
* 122164 122263: contig of 20443 bp in length
* 122264 143900: gap of unknown length
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    chromosome       "16"
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BASE COUNT 37791 a 34128 c 34320 g 35886 t 1775 others
 ORIGIN

Query Match 10.1%; Score 208; DB 2; Length 143900;
 Best Local Similarity 100.0%; Pred. No. 4e-98;
 Matches 208; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1861 agagacataatggaagatgaatgacttcttgaagaagcgaatgcccacaaatata 1920
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QY 1921 ccttgattggagatcaccaacgaatccttgaacagcatctcgaatcccttcaagtag 1980
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DB 27278 CCGTGATTGGGATCACACCAAGCTCTTTACACGCAATTTCCAGACTCTTTACAGTAGTG 27219
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QY 1981 tggatctccaccccgatgltgacatgaaccacgactccctctacacgcagaataattgtga 2040
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DB 27218 TGAGCTCCGCCACCGCTGTGTGACATGCAAGCCAGTCCCTCTGACGGCACAATTTGTCA 27159
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QY 2041 ctgagatgtagacttggagattcccat 2068
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DB 27158 CTGAGATGATGACATTGGCATTCGCCAT 27131
  
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RESULT 10

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 LOCUS 175691 bp DNA linear HTG 03 JUL 2001
 DEFINITION Homo sapiens chromosome 16 clone RP11-528K16, WORKING DRAFT
 SEQUENCE, 1 ordered pieces.

AC007615
 AC007615.6 GI:14589428
 HTG: HTGS_PHASE2: HTGS_DRAFT; HTGS_ACTIVEFIN.
 SOURCE
 ORGANISM
 Homo sapiens

REFERENCE
 AUTHORS
 TITLE
 JOURNAL
 REFERENCE
 AUTHORS
 1 (bases 1 to 175691)
 2 (bases 1 to 175691)
 Bruce, D., Mundt, M., Douget, N., Monk, C., Saunders, E., Robinson, J., Jones, M., Buckindham, J., Chastock, L., Thompson, S., Goshwin, J., Bryant, J., Fawcett, J., Melnick, L., Langmuir, J., White, S., Tetam, J., Campbell, C., Fawcett, J., Malthe, M., Russell, M., Sutherland, R., McKerry, K., Han, C. and Heaven, L.
 Direct Submission
 Submitted (20-MAY-1999) center for Human Genome Studies, DOE Joint Genome Institute, Los Alamos National Laboratory, MS M888, Los Alamos, NM 87545, USA
 On Jul 3, 2001 this sequence version replaced gi:13928651.

COMMENT
 JOURNAL
 TITLE
 JOURNAL
 REFERENCE
 AUTHORS

Sequence Quality Assessment:
 This entry has been annotated with sequence quality estimates computed by the Phrap assembly program.
 All manually edited bases have been reduced to quality zero.
 Quality levels above 40 are expected to have less than 1 error in 10,000 bp.
 Base-by-base quality values are not generally visible from the GenBank flat file format but are available as part of this entry's ASN.1 file.

Sequence Quality Assessment:
 This entry has been annotated with sequence quality estimates computed by the Phrap assembly program.
 All manually edited bases have been reduced to quality zero.
 Quality levels above 40 are expected to have less than 1 error in 10,000 bp.
 Base-by-base quality values are not generally visible from the GenBank flat file format but are available as part of this entry's ASN.1 file.

Sequence Quality Assessment:
 This entry has been annotated with sequence quality estimates computed by the Phrap assembly program.
 All manually edited bases have been reduced to quality zero.
 Quality levels above 40 are expected to have less than 1 error in 10,000 bp.
 Base-by-base quality values are not generally visible from the GenBank flat file format but are available as part of this entry's ASN.1 file.

FEATURES
 source
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 organism "Homo sapiens"
 db_xref "taxon:9606"
 chromosome "16"
 clone_name "RP11-528K16"
 clone_lib "RP11 human BAC library 11"

BASE COUNT 42392 a 42125 c 45277 g 45896 t 1 others
 ORIGIN

Query Match 9.0%; Score 187; DB 2; Length 175691;
 Best Local Similarity 100.0%; Pred. No. 4.7e-87;
 Matches 187; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB 31036 TACTGTAAATCATGCTAGATCTTTGCTTAAGCTGTCACATATATTTAATAACTA 31065
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001175(1997)0441(17)

TITLE	JOURNAL	REFERENCE	AUTHOR'S TITLE	COMMENT
Direct Submission	Submitted (22-AUG-1997)	The Institute for Genomic Research, 9712 Medical Center Dr., Rockville, MD 20850, USA	3 (bases 1 to 262004)	
ADAMS, M.D. and Lettner, B.J.				
Direct Submission	Submitted (28-JUL-1998)	The Institute for Genomic Research, 9712 Medical Center Dr., Rockville, MD 20850, USA		
On Jul 31, 1998 this sequence version replaced gi:2440964.				
BAC clone, C119815K.270G1 is located in chromosome 16. Genes were identified by a combination of five methods: XORAT (available by anonymous ftp from arthur.epm.ornl.gov), GeneIndex (available by anonymous ftp from colin@u.washington.edu), GENSCAN (available using the e-mail server at genescanomic.stanford.edu), searches of the EST database at tigr (http://www.tigr.org/tsh/hcd/hcd.htm) and searches against a peptide database. Reports were identified using RepeatMasker (Smith, A. and Green, P., unpublished, http://ftp.genome.washington.edu/rm/RepeatMasker.html).				
Location/Qualifiers				
1. .202004				
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/protein_id="AAC27823.1"				

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Best Local Similarity	100.00%;	Prod. No. 3e-81;	
Matches 176;	Conservative 0;	Mismatches 0;	Indels 0; Gaps 0;
STS	STS	STS	STS
BASE COUNT	56977 a 45624 c 44447 g 54956 t		
ORIGIN			

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10 66278 caaaagcaatcttgaatgttgaatctccatggagcaatcttactcagactgacgaagctgaat 1458
07 1459 tctccagacttcttcttcttcttcttcttcttcttcttcttcttcttcttcttcttcttctt 1514
10 66308 tctccagacttcttcttcttcttcttcttcttcttcttcttcttcttcttcttcttcttctt 1514

Search completed: July 30, 2002, 05:44:49
Job Time: 8689 sec


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SOURCE          human.
ORGANISM        Homo sapiens
Eukaryotic: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi:
Mammalia: Eutheria: Primates: Catarrhini: Homiidae: Homo.
REFERENCE      1 (bases 1 to 788)
AUTHORS        NIH-MGC http://mgs.nci.nih.gov/.
TITLE          National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL        Unpublished (1999)
COMMENT        Contact: Robert Strausberg, Ph.D.
                Email: rcstra@mail.nih.gov
                Tissue Procurement: ATCC
                cDNA Library Preparation: CLONTECH Laboratories, Inc.
                DNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
                DNA Sequencing by: Incyte Genomics, Inc.
                Clone distribution: MGC clone distribution information can be
                found through the I.M.A.G.E. Consortium/LNL at:
                http://image.llnl.gov
                Plate: LDCM451 row: 1 column: 08
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                        /clone_1lb="NIH_MGC_61"
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                        /lab_host="DH10B (T1 phage-resistant)"
                        /note="Organ: testis; Vector: pDRM-11B (Clontech); Site:
                        Still (qqccctcgcgc); Site:2: Still (qqccatatagc);
                        Double-stranded cDNA was prepared from cell line RNA. 5'
                        and 3' adaptors were used in cloning as follows: 5'
                        adaptor sequence: 5'-CACGCCCATATGCGC-3' and 3' adaptor
                        sequence: 5'-ATTCTAGACGCCGACGCGCCGACATG-dT(30)BN-3'
                        (where B = A, C, or G and N = A, C, G, or T). Average
                        insert size 1.75 kb (range 0.9-4.0 kb). 15/15 colonies
                        contained inserts by PCR. This library was enriched for
                        full-length clones and was constructed by Clontech
                        Laboratories (Palo Alto, CA). Note: this is a NIH_MGC
                        Library."
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alignment_block:
US-09-932-678-2 x BC502871 ..
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2 CGGCTCGACGAGATGTCATCTCTACGCCGACGCCCGACATGCAATTCAT 51
|||||
318 TIsertLeuValIleuSerTyrMetLeuAspAlaCysTyrValAspGlyIysV 335
|||||
52 GCTCTTGCTTTGCTCTACATGAGGATGTCCTCTATGAGTGGTGGTAAAGG 101
|||||
335 aIAspAsnGlyIysThrIysAspLeuTyrArgAspLeuIleasnIlePhe 351
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102 TTGATTAACGGCAAAACAAAGAGATATATACGGACCTATAACATCTTT 151
|||||
352 AspIysLeuLeuLeuProThrHisAlaSerCysHisValGlnPhePhe 368
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152 GACAACCTCTGTTGCCACCATGCTCTCCATGTCATACATTTTCAT 201
|||||
368 tPheTyrLeuGlySerPheIysLeuGlyPheAlaGluAlaPheLeuGluH 385
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seq_name: jh_est2.BF203684

seq. documentation link: [seq. documentation link](#)

LOCUS BF203684 996 bp mRNA linear EST 06-NOV-2000

DEFINITION 601866331P1 NIH_MGC_17 Homo sapiens cDNA clone IMAGE:4093348 5'

ACCESSION BF203684

VERSION BF203684.1 GI:11097270

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 996)

AUTHORS NIH-MGC <http://mgi.nhl.nih.gov/>.

TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL Unpublished (1999)

COMMENT Contact: Robert Strausberg, Ph.D.
Email: cepb@small.nhl.gov
Tissue Procurement: ATCC
cDNA Library Preparation: Ling Hong/Kubin laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LMNI)
DNA sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LMNI at: image.llnl.gov
Plate: LCM967 row: a column: 07
High quality sequence stop: 684.
Location/Qualifiers

1..996


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17 rSerSerAlaValIleIleLeuGlyAlaSerArgThrGlyIleSerAsn 34
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73 GTCTCTGCAGATTAAAGAGCTGGGCGCTGGAGACATGGCATTTCAATA 122
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67 uThrAspPheGluLeuLeuLysAsnGluLeuLeuAspProAspIleL 84
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184 sPThrCysHisArgAlaLeuGlnIleIleAlaArgTyrValProSerThr 200
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  ACCESSION BG428305
  VERSION BG428305.1 GI:13334811
  KEYWORDS EST.
  SOURCE human.
  ORGANISM Homo sapiens
  Eukaryota; Metazoa; Chordata; Granulata; Vertebrata; Euteleostomi;
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  REFERENCE 1 (bases 1 to 1037)
  AUTHORS NIH-MGC, http://mml.nhl.nih.gov/.
  TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

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JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
  Email: rchap3s@mail.nih.gov
  Issue Procurement: Clontech Laboratories, Inc.
  cDNA Library Preparation: Clontech Laboratories, Inc.
  DNA Sequencing by: Incyte Genomics, Inc.
  Clone distribution: MGC clone distribution information can be
  found through the I.M.A.G.E. Consortium/ILN at:
  http://image.llnl.gov
  Plate: LTM1361 row: e column: 22
  High quality sequence stop: 637.
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  sequence: 5'-CACGGCATATGACC-3' and 3' adaptor sequence:
  5'-ATCTAGAGCGCGGAGCGGCGACATG-dl(30)BN-4' (where B = A,
  C, or G and N = A, C, G, or T). Average insert size 1.65
  kb (range 0.5-4.0 kb). 15/15 colonies contained inserts
  by PCR. This library was enriched for full-length clones
  and was constructed by Clontech Laboratories (Palo Alto,
  CA). Note: This is a NIH-MGC library."
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    |||
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  173 CGGTTTGAGTGAAGCTGGACAAAGTCTTCTGAGTACAAAAAGGCTGA 222
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  84 yAspAspGlnIleIleAsnTrpLeuLeuGluPheArgSerSerIleMet 100
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  273 AGCATGACCAATCATCAACTGGTGTCTAGAAATTCGGTCTTCAATCATG 321
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  101 TyrLeuThrLysAspPheGluGluLeuIleSerIleIleLeuArgLeu 117
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http://image.jhu.edu
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 Location/Qualifiers

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/clone "IMAGE:5268011"

/clone_lib "NIH_MGC_97"

/lab_host "DH10B"

/note "organism: pooled colon, kidney, stomach; Vector: pMVS-Spacer; Site 1: Not; Site 2: Eovk (destroyed); RNA

source: anonymous pool of 3 colonies, age 26 yo male, 49 yo female, 71 yo male colon; 46 yo female kidney, and pool of 2 stomachs; 62 yo male and 70 yo female. Library is

oligo dt primed and directionally cloned (Eovk site is destroyed upon cloning). Average insert size 1.4 kb,

insert size range 1-3 kb. Library is normalized and

enriched for full-length clones and was constructed by C. Gabor (Invitrogen). Research Genetics tracking code

BASE COUNT 203 a 215 c 207 g 203 t
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alignment sources:

quality: 1000,50 length: 225
 Ref ID: 4,904 gaps: 4
 Percent Similarity: 99.667 Percent Identity: 87.111

alignment block:

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 203 GATGATGCAAGTAAATCTGCAATGAGCAAGGATGCGTCAAAATATA 252
 407 ccGlyscPhclagclAlagclAlagclshclclclclclclclcl 423
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 273 TTTGCAAGTCTTTGCAAGCAAGCAAAATTTATGCTTATCTGTA 402
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603 TTTGCTAAAGATTCGCAAGCAAGTAAATCTGCAAGTCTTGCAAT 652
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 573 cl 753
 753 GCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 802
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seq_documentation_block:

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Accession: M1464721 mRNA sequence.

Version: M1464721.1 GI:15255464

KeyWords: EST.

Source: human.

Organism: Homo sapiens.

Eukaryotic; Metazoa; Chordata; Cladoda; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo;

1 (bases 1 to 746)

NIH-MGC hybrid image: m1464721.

Title: NIH-MGC hybrid image: m1464721.

Journal: National Institutes of Health, Mammalian Gene Collection (MGC)

Comment: Deposited (1999)

Contact: Robert Stansberg, Ph.D.

Email: eqapus@mail.nih.gov

Tissue Procurement: Miklos Palkovits, M.D., Ph.D.

cDNA Library Preparation: Michael J. Brownstein (NIH/RI), Shitaki

Toshioyuki and Piero Carninci (RIKEN)

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LMNI)

RNA Sequencing by: Invitrogen, Inc.

Clone Distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/IMB at:

http://image.jhu.edu

Plate: UAM1459, row: b, column: 12

High quality sequence stop: 725.

Location/Qualifiers

1..746

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/db_xref "taxon:9606"

/clone "IMAGE:5268011"

/clone_lib "NIH_MGC_97"

/lab_host "DH10B"

/note "organism: testis; Vector: pBluescript (modified

phagescript KS+); Site 1: Hamby site 2: Salt Hole (GTCGAG

) oligo-dt primed using primer 5'-ATTGTTTCTTTTATVN-3',

size-selected for average insert size 2.2 kb and

normalized to 10⁶ 5'. This is a primary library enriched

for full-length clones and constructed using the

cap-trapper method (Guarnieri, in preparation). Library

constructed by M. Brownstein (NIH/NIHRI, National

Institutes of Health). Note: This is a NIH_MGC library."

BASE COUNT

206 a 143 c 178 g 209 t

ORIGIN

alignment sources:
 quality: 989,50 length: 234
 Ref ID: 4,477 gaps: 4
 Percent Similarity: 94.444 Percent Identity: 89.744
 alignment_block:

US-09-932-678-2 x B1464721

Align seq 1/1 to: B1464721 from: 1 to: 736

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141 TGGCTGATTAGAGAAAGACTTTTTCATTTCTCCCAAGAAATCGTT 190
51 ArgPheGlyGlyThrValThrGluValLeuLeuLysTyr..LysLysGly 66
191 CGGTTTGGTGGAACTGTCACACGAGCTTCTGAGTACACACACACGCT 240
67 GluThrAsnAspPheGluLeuLeuLysAsnGluLeuLeuAspProasp 83
241 GAACAACATGACTTTCAGTTGTTGAAGAACACGCTGTTAGATCCACACAT 290
83 eLysAspAspGluLeuLeuLeuLeuLeuGluProArgSerSerLem 100
291 AAGAGATGACCAAGATCATCTACCTGGCTGAGAACTGCTTCTTATCA 340
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341 TGACTTGGACAAAAGACTTGGACCACTTATTCAGTATATTAATCAATG 390
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541 TGTAAATGTTTCAGATTCATGATGATCAAGATGATATCTTCGATCGAA 590
183 Phe..AspThrCysHisArgAlaLeuGlnLleLleAlaArgTyrVal 198
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198 oSerThrProTrpPheLeuMetProLleLeuValGluLysPheProPhe 215
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seq_documentation_block: 563 bp mRNA linear EST 16-FEB-2000

LOCUS AM408066

DEFINITION U1-HF-BM0-adu-b-10-0-U1.N1H.MGC_38 Homo sapiens cDNA clone

IMAGE: 3062754 5', mRNA sequence.

ACCESSION AM408066

VERSION AM408066.1 GI:6927123

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 563)

AUTHORS

TITLE NIH-MGC http://mgc.nhl.nih.gov/.

NATIONAL

INSTITUTES

OF HEALTH.

MAMMALIAN

GENE

COLLECTION (MGC)

JOURNAL.

UNPUBLISHED

(1999)

CONTACT:

ROBERT

STRAUSBERG,

PH.D.

EMAIL:

cgsosr@mail.nih.gov

EVO R1

SITE

SHOWN

AT

THE

BEGINNING

OF

THE

SEQUENCE.

TISSUE

PROCUREMENT:

LOUIS

M. STAUDT,

M.D.,

PH.D.

CDNA

LIBRARY

PREPARATION:

M.B. SOARES

LAB

DNA

SEQUENCING

BY:

M.B. SOARES

LAB

CLONE

DISTRIBUTION:

MGC

CLONE

DISTRIBUTION

INFORMATION

CAN

BE

FOUND

THROUGH

THE

L.M.A.G.E.

CONSORTIUM/HLNI

AT:

WWW.BIO.NHL.GOV/BLHP/IMAGE/IMAGE.HTML

Seq primer: M13 forward.

FEATURES

Source

Location/Qualifiers

1..563

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:3062754"

/clone="U1-HF-MGC_38"

/issue="type="lymph"

/cell_type="germinal center B cells"

/lab_host="DH10B (RT)"

/note="Vector: pTZ19-Pac; Site 1: NotI; Site 2: Eco RI;

Constructed from size fractionated cytoplasmic mRNA

(2.5-3.5kb). Directionally cloned. Cells provided by Louis

M. Staudt, Ph.D. Library preparation by Maria de Fatima

Ronaldo, Ph.D. and M. Bento Soares, Ph.D."

BASE COUNT

151 a 122 c 119 g 171 t

ORIGIN

alignment_scores:

Quality: 966.00 Length: 185

Ratio: 5.250 Gaps: 1

Percent Similarity: 99.459 Percent Identity: 99.459

alignment_block:

US-09-932-678-2 x AM408066

Align seq 1/1 to: AM408066 from: 1 to: 563

330 TyrValAspGlyIysValAspAsnGlyLysThrArgLeuArgAs 446

10 TATGTAGATGATAGCTGATAGGCAAAACAAAGATGATATATGCGCA 59

346 pCulleAsnLlePheAspLysLeuLeuLeuLeuLeuLeuLeuLeuLeu 464

60 CTGATTAACATCTTGAACAAACTCTGTTGTCACCCATGCTGCTGCTG 109

363 LysValGlnPhePheMetPheTyrLysCysSerPheLysLeuGlyPheVal 379

110 ATGTACAGGTTTTCATGCTTTTACCTCTGTAGTTTCAATATGATATGCA 159

380 GluAlaPheLeuGluHisLeuTrpLysLysLeuGluAspProSerAsp 396

160 GAGGCAATTTTGAACATCTCTGAAAAAATTCAGAGACCAAGTATATCC 209

396 GAlaLleLeuArgGlnAlaAlaGlyAsnTyrTleGlySerPheLeuAla 414

210 TCCATCATCATGACAGAGCTGCTGGAAATTAATTTGGAAATTTTGGTAA 259

413 rAlaLysPheLleProLeuLleThrValLysSerTyrLysLeuLeuLeu 429

260 GAGCTAAATTTATTCCTCTTATTTACTGTAAATCATGCTTAGATCTTTTG 309

430 ValAsnTrpLeuHisLleTyrLeuAsnAsnGluAspSerGlyThrLysAl 446

310 GTTAATGGCTGCAATATACCTTAATTAAGCAGGATTCGGGAAACAAAGT 459

446 aPheCysAspValAlaLeuHisGlyProPheTyrSerAlaCysGlnAlaVal 464

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|||||
400 ATGTTGATGCTTCTCTGATGACCAATTTATCTAGCTCTGCAAGCTG 409
403 aTPhcValPheValPheArgHisLysGlnGlnGlnGlnGlnGlnGln 479
|||||
410 TGTTCACACCTCTTGCTTTACAGACACAGACAGCTTTTGAGCGGAAACCTG 459
|||||
400 TgcStGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGln 496
|||||
460 AAATAAAGCTCTTCTAGCTCTGAAATTTTGACGCAATATGTA 509
|||||
496 GATGCTGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGln 512
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510 TCACTAGCTAAATCTCTGAAATTTGCTCTGCTGCTTAACTT 559
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513 Phe 513
|||||
560 TTT 562

seq_name: gb_est.2:H549394
seq_documentation_block:
LOCUS B1649394 727 bp mRNA linear EST 05-SEP-2001
DEFINITION G0439004991 NIH_MGC_75 Homo sapiens cDNA clone IMAGE:5261100 5'.
ACCESSION B1649394
VERSION B1649394.1 GI:15446706
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catartida; Homnidae; Homo.
REFERENCE 1 (bases 1 to 727)
AUTHORS NIH_MGC http://mgl.nhl.nih.gov/
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: eqdphs@mail.nih.gov
Tissue preparation: Miklos Pakovits, M.D., Ph.D.
cDNA library preparation: Michael J. Brownstein (NIH), Shiroki
Toshitoki and Piero Carninci (RIKEN)
cDNA library arrayed by: The J.M.A.G.E. Consortium (JLM)
DNA sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the J.M.A.G.E. Consortium/JLM at:
http://image.jlm.gov
plate: JLM11658 row: b column: 14
high quality sequence: slope: 715.
FEATURES
Source
1..727
Annotation "Homo sapiens"
db_xref "taxon:9606"
/codon "IMAGE:5261100"
/codon "lib NIH_MGC_95"
/issue type "HaploCampus"
/lab host "H10B"
note "Organic brain; Vector: pBluescriptR (modified
phoscript KS2); Site 1: BamHI; Site 2: SalI-XbaI (q1eqn
); clone id primed using primer 5'-TTTTTTTTTTTTTTT-VN-47',
size selected for average insert size 2.5 kb and
normalized to rot 5. This is a primary library enriched
for full length clones and constructed using the
cap trapped method (Carninci, in preparation). Library
constructed by M. Brownstein (NIH/NCRI, National
Institutes of Health). Note: This is a NIH_MGC Library."
BASE COUNT
Hz a 191 c 171 g 181 t
alignent sources:
quality: 966.00 length: 197
ambigu: 5.031 gaps: 1
percent similarity: 97.462 percent identity: 94.401

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alignent_block:
us-09-932-678-2 x B1549394
Align seq 1/1 to: B1549394 from: 1 to: 727
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|||||
98 TATCGACAGAGATTCGACAGACATTTTGGAAATCTTGCAGAAAT 147
|||||
490 GATGAGGATGCAAGTATCTGCTGATCATCAGCGAGATGCTGAAATATA 197
|||||
148 GATGAGGATGCAAGTATCTGCTGATCATCAGCGAGATGCTGAAATATA 197
|||||
407 TgcLysGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGln 423
|||||
198 TGGAGAGCTTTTGGCAGACGTAATTTATTCGCTTATTA-TGTAATA 247
|||||
424 SGTGysLacAspLeuLeuValAsnTrpLeuHisLeuTyrLeuAsnAsp 440
|||||
248 CATGCTCATATCTTTGGTAACTGCTGCAATATACCTTAATATATTA 297
|||||
440 GASPSTGCTGlyTPhLysAlaPheGysAspValAlaGlnHisGlyProPhe 457
|||||
298 GATTCGAGACCAAGCAATCTGCAATCTGCTGCTGCTGCTGCTGCTG 447
|||||
457 YTSGLAcysGlnAlaValPheTyrPheValPheArgHisLysGln 473
|||||
348 ATTGACAGCTGCGACAGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 497
|||||
474 TgcLysGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGlnGln 490
|||||
398 GTTTCGAGCAAAATCTGAAAGAAATTTTGAGATCTGCAAGCTGTGAA 447
|||||
490 GSPSTGATGlyLeuAlaMetSerGlnGlnGlnGlnGlnGlnGlnGlnGln 507
|||||
448 TTTTCGCGCATAGTATGATGATGATGATGATGATGATGATGATGATG 497
|||||
507 TSGValValAsnPhoPheAlaAlaLeuThrAspLysTyrLeuVal 523
|||||
498 GTTCATGCTGTTAGCTTTTATCTGCAATTCACAAATATATGACGCTGC 547
|||||
524 PgcTyrTyrThrLeuLeuAlaGlnAsnAsnAcGlnMetLeuProVal 540
|||||
548 TGTGCTGACATCATCATGACAGCAAGCAATCCGCAATGCTGCTCATAT 597
|||||
546 GATGAGGATGATGAGGATGATGATGATGATGATGATGATGATGATG 557
|||||
554 TACGACTGCAATCTGACGACATGATGATGATGATGATGATGATGATG 647
|||||
554 SGTPhcPheProPheAspProGysValLeuLysAsn 569
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648 AATGCTTAACTGCTTTCATCTGCTGCTGCTGCTGCTGCTGCTGCTG 606
|||||

seq_name: gb_est.2:B6167061
seq_documentation_block:
LOCUS B6167061 990 bp mRNA linear EST 06-FEB-2001
DEFINITION G0244474091 NIH_MGC_89 Homo sapiens cDNA clone IMAGE:445509 5'.
ACCESSION B6167061
VERSION B6167061.1 GI:12674764
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catartida; Homnidae; Homo.
REFERENCE 1 (bases 1 to 990)
AUTHORS NIH_MGC http://mgl.nhl.nih.gov/
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: eqdphs@mail.nih.gov

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Tissue Procurement: ATCC
 cDNA Library Preparation: Life Technologies, Inc.
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/ILMI at:
<http://image.llnl.gov>
 plate: L1AM10247 row: 0 column: 10
 High quality sequence stop: 665.
 Location/Qualifiers
 source
 1..990

/organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:4455009"
 /clone_lib="NIH_MGC_89"
 /issue_type="hypermethylation, cell line"
 /lab_host="DH10B (phage-resistant)"
 /note="Organ: Kidney; Vector: pCMV-SPORT6; Site 1: Not;
 Site 2: Salt; Cloned unidirectionally; oligo-dT primed.
 Average insert size 1.3 kb. Library enriched for
 full-length clones and constructed by Life Technologies.
 Note: This is a NIH_MGC library."
 BASE COUNT 304 a 275 c 219 g 192 t
 ORIGIN

alignment_scores:
 Quality: 935.00 Length: 187
 Ratio: 5.109 Gaps: 1
 Percent Similarity: 97.861 Percent Identity: 97.326

alignment_block:

US-09-932-678-2 x BG167061 ..

Align seq 1/1 to: BG167061 from: 1 to: 990

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 2 ACCTTTGTTTTAGACACAGCAGCTTTTGACCGAAGACCTGAAGAG 51
 482 YLeuGlnTyrLeuGlnSerLeuAsnPheluarqLValMetSerL 499
 |||||||
 52 TTTCAGATCTTCAGAGTCGATTTGAGCGATAGTATGACGACG 101
 499 eu AsnProLeuLysLLeuGlnProSerValValAsnPhelua 515
 |||||||
 102 TAAAGTCCCTGAGAGATTTCCTCCCTCAAGTCTTAACCTTTT 151
 515 alleThrAsnLysTyrGlnLeuValPheCysTyrThrLleGln 532
 |||||||
 152 AATACAAATAGTACGACGCTGCTTCTGCTACACCATCATTA 201
 532 suAsnATGlnMetLeuProValLleArqSerThrAlaGlyLys 548
 |||||||
 202 ACATTCGCCAGATCTCCGACATCATAGAGTACCCGTGAGGAG 251
 549 ValGlnLleYstrAsnProLeuAspThrPhePheProPheAs 565
 |||||||
 252 GTGACGATCTGACCAACCCGCTGGACACCTTCCCTTGGAT 301
 565 sValLeuLysArqSerLysLysPheLleAspProLleTyrGln 582
 |||||||
 302 TGCTGTAAAGCTCAAGAAATTCATGATCATTTATAGGTGG 351
 582 LuAspMetSerAlaGlnLleuGlnLleuPheLysLysPromet 598
 |||||||
 352 AAGCATGACGCTGACAGAGCTACAGAGTTCAGAAACCATGA 401
 599 AspLleValGluAspGluAspAspPheLeuLysGlnValPro 615
 |||||||
 402 GACATAGTGAAGATGAAGATGATGCTTCTGAAGACGAGGAG 451
 615 uAsnAspThrValLleGlyLleThrProSerSerPheAspThr 632

|||||
 452 GAATGATAGCTGATGATGATGATGATGATGATGATGATGAT 501
 632 rSerProSerSerValGlySerProValLeuTyrMetuLPro 648
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 502 GAAGTCTTCAAGTATGAGTGGCTGGCAAGCTGTAGTACAGCA 551
 649 SerProLeu 651
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 552 ATTCCTCTC 560

seq_name: gb_est2:BG297178

seq_documentation_block:

LOCUS BG297178 1045 bp mRNA linear EST 21 FEB 2001
 DEFINITION 602395174P1 NIH_MGC_94 Mus musculus cDNA clone IMAGE:4506594 5,
 mRNA sequence.

ACCESSION BG297178

VERSION BG297178.1 GI:13060570

KEYWORDS EST.

SOURCE house mouse.

ORGANISM

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Scurionathia; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 1045)

AUTHORS NIH-MGC <http://mgs.cni.nih.gov/>.

TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

UNPUBLISHED (1999)

CONTACT: Robert Strausberg, Ph.D.

EMAIL: gaasb@mail.nih.gov

Tissue Procurement: The Coriell Laboratory

cDNA Library Preparation: Life Technologies, Inc.

DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/ILMI at:

<http://image.llnl.gov>

plate: L1AM10382 row: d column: 19

High quality sequence stop: 687.

FEATURES

source

1..1045

/organism="Mus musculus"

/db_xref="taxon:10090"

/clone="IMAGE:4506594"

/clone_lib="NIH_MGC_94"

/issue_type="Fetal"

/lab_host="DH10B (phage-resistant)"

Site 2: Salt; Cloned unidirectionally; oligo-dT primed.

Average insert size 3.3 kb. Library enriched for

full-length clones and constructed by Life Technologies.

Note: This is a NIH_MGC library."

BASE COUNT

303 a 225 c 273 g 244 t

1 others

alignment_scores:

Quality: 928.50 Length: 336

Ratio: 3.613 Gaps: 10

Percent Similarity: 76.488 Percent Identity: 64.881

alignment_block:

US-09-932-678-2 x BG297178 ..

Align seq 1/1 to: BG297178 from: 1 to: 1045

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 26 ATGGGAGGCGGAGTCTTCACAGCGCTTCTCGGCGATGACGAT 78
 17 pSerSerAlaValLysLysLysValAspArqPheGlyLysGAs 44
 |||||||
 76 GAGCTCTGATACG.....CTGAGTGGGTCGGAGAGTGGGCTT 122


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303 TTGAGCAAGCTATCAGTATTATTAAGATTGCTTGGTCAATAGAG 352
122 TGTGTTGValValGUGUGUtyrLeuAlaPheLeuGlyAsnLeuValSerA 139
|||||
353 TCAACAGTAGTGGAGAGATATTGGCTTTCTTGTAATCTTGATCAG 402
139 TGGTThrValPheLeuArgProCysLeuSerMetIleLeuSerHisPhe 155
|||||
403 CACAGACTTTTCTCCAGAGCCGTGCTTACAGATGATCTCTCCATTTT 452
156 ValProPheArgValIleIleLeuLysGluGlyAspValAspValSerAsp 172
|||||
454 GAGCTCCCGGAGTATCATTAAGCAAGCCGATGTAGATGTTTCAGATT 502
503 CTCATGATGATACGATGATGATATCTCTCTGCANATTTCACCATGTCCACA 551
189 AlalPheGlnIle 192
|||||
552 GCCCTGCCAATA 563
seq_name: qb_est2:BC295610
seq_documentation_block:
LOCUS BC295610 956 bp mRNA linear EST 21-FEB-2001
DEFINITION 60239299.2P1 NIH_MGC_94 Mus musculus cDNA clone IMAGE:4504751 5',
ACCESSION BC295610
VERSION BC295610 GI:13057417
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 956)
AUTHORS NIH-MGC http://mgc.ncbi.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: rgsb@fmail.nih.gov
Tissue Procurement: The Cepko Laboratory
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: L1AM10377 row: q column: 24
High quality sequence stop: 660.
location/Qualifiers
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/organism="Mus musculus"
/db_xref="taxon:10090"
/clone_image="IMAGE:4504751"
/clone_id="NIH_MGC_94"
/tissue_type="retina"
/lab_host="DH10B (phage-resistant)"
/notes="Organ: eye; Vector: pCMV-SpOK16; Site_1: NotI;
Site_2: SalI; Cloned unidirectionally; oligo-dT primed.
Average insert size 3.3 kb. Library enriched for
full-length clones and constructed by Life Technologies.
Note: This is a NIH-MGC library."
BASE COUNT 266 a 204 c 239 g 247 t
ORIGIN
alignment_scores:
Quality: 898.50 Length: 277
Ratio: 3.682 Gaps: 8
Percent Similarity: 88.087 Percent Identity: 71.841
alignment_block:

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US-09-932-678-2 x RG295610
Align Seq 1/1 to: HG295610 from: 1 to: 956

147 CysineSerMet1Lea1aSerHisphenValProPrArGVal11e1lely 163
||||| ||||||| ||||||| ||||||| ||||||| |||||||
4 TGTGTGCAATGATGGTGTCAATTGTACCTGGCGAGTAATGTCAA 54
||||| ||||||| ||||||| ||||||| ||||||| |||||||
163 sGluGlyAspValAspValSer AspSerAspaspGluAspaspGlu 179
||||| ||||||| ||||||| ||||||| ||||||| |||||||
52 GCAAGGTGATATATATATTTACCATTTGATGACCAAGATGCAATGA 101
||||| ||||||| ||||||| ||||||| ||||||| |||||||
180 prAla1aSerPhe AspThrCysHisArGAlaGlu11e1Le1aArG 196
||||| ||||||| ||||||| ||||||| ||||||| |||||||
102 TCGTGCAATTTTGACACATGTCCACAGAGCTTGCAATATATACAAAT 151
||||| ||||||| ||||||| ||||||| ||||||| |||||||
156 yValProSerThrProThrPheLeuMetPro11eValu1GluysPhe 212
||||| ||||||| ||||||| ||||||| ||||||| |||||||
152 ATATCCCATTCACACATCGTTTCTATATGCCAATCTGGTGAGAAATTC 201
||||| ||||||| ||||||| ||||||| ||||||| |||||||
213 prPheValArG1ysSerG1uArvThrGlu1GcysTyValHisAsnG 229
||||| ||||||| ||||||| ||||||| ||||||| |||||||
202 CCGTTGTGAGCAATGCCAGACAAATTGAAATGTAATTTATACATT 250
||||| ||||||| ||||||| ||||||| ||||||| |||||||
229 uLeuArG1eSerVal1yrPheProThrLeuArG11eGlu1e1eGlu1e 246
||||| ||||||| ||||||| ||||||| ||||||| |||||||
251 ATTAAGATATAGTTTATATTTCCCACTTTGAGCGCGTAATTTGAGAG 300
||||| ||||||| ||||||| ||||||| ||||||| |||||||
246 eAl1e1eGlu1ysLeuLeu1ys1euaSpValAsnAlaSerArG1u1y 262
||||| ||||||| ||||||| ||||||| ||||||| |||||||
301 T1G1ATTTGAAAGCTACTACCTATGATGTGATGTGGCGAGAT 350
||||| ||||||| ||||||| ||||||| ||||||| |||||||
263 T1eGluSpAlaGlu1uThrAlaThrG1uThrCysG1y1yThrAspSe 279
||||| ||||||| ||||||| ||||||| ||||||| |||||||
351 ATTAACATGTGTGAGGAAGCAAGCACTTGCTGTGGCACAATAC 400
||||| ||||||| ||||||| ||||||| ||||||| |||||||
279 rThrGlu1yLeuPheAsnMetAspG1uAspG1uGlu1ThrGluHis1u 296
||||| ||||||| ||||||| ||||||| ||||||| |||||||
401 CCGAGAGCACTGTTATATATGATGAGATGAGGACACTGACCTGACA 450
||||| ||||||| ||||||| ||||||| ||||||| |||||||
296 br1ys1aG1yProG1uArG1euaSpG1uMetValHisProVal1aG1u 312
||||| ||||||| ||||||| ||||||| ||||||| |||||||
451 ATAAAGCTGATACAGACAGAGCTTAACAGATGGTCACTGACCGAGAG 500
||||| ||||||| ||||||| ||||||| ||||||| |||||||
313 ArG1euaSp11eLeuMetSer1euaVal1euaSer1yMet1ysAspVal1y 329
||||| ||||||| ||||||| ||||||| ||||||| |||||||
501 C1eCTGCACTGCTGCTGTGCTGTGCTGTGCTGCTGCTGCTGCTGCTG 550
||||| ||||||| ||||||| ||||||| ||||||| |||||||
329 sTyValAspThr1ysValAspaspG1y1y1yThr1ysAsp1e1y1rArA 346
||||| ||||||| ||||||| ||||||| ||||||| |||||||
551 CCGTGTACACCTTAATAATATATACCAATAAACCAAGATTATATCGTG 600
||||| ||||||| ||||||| ||||||| ||||||| |||||||
346 sPhe11eAsn11ePheAsp1y1eua...1eua1eProThrHisAlaSe 361
||||| ||||||| ||||||| ||||||| ||||||| |||||||
601 ATGTGATATGATCTTTGTGAATAAAGCTTTGTTGTGGCCACAAATGCTC 650
||||| ||||||| ||||||| ||||||| ||||||| |||||||
361 rCysHisVal1eInPhe...MetPheTy1euaCysSerPhe1ys1euaG 377
||||| ||||||| ||||||| ||||||| ||||||| |||||||
651 CTGGCATGTACCGTCTTTCATCTTTTCTGCTGGCAGAGCTTCACTGG 700
||||| ||||||| ||||||| ||||||| ||||||| |||||||
377 1yPheArG1a1aPhe1eua1y1eua1y1y1y1y1y1y1y1y1y1y1y 394
||||| ||||||| ||||||| ||||||| ||||||| |||||||
701 GATATGCCAGAGCAATTTTGACATCTCTGGGACCAAGTACCGATCCA 750
||||| ||||||| ||||||| ||||||| ||||||| |||||||
394 sArAsnPrAla11e1eArG1u1a1a1a1a1a1yAsnTy11e1ySerP 410
||||| ||||||| ||||||| ||||||| ||||||| |||||||
751 ATACCC...CCCATATACGAGAG...CGGTGCAAAATATTTGGAGAGCT 794
||||| ||||||| ||||||| ||||||| ||||||| |||||||
411 e1eua1a1a1a1a1y1y1y1y1y1y1y1y1y1y1y1y1y1y1y1y 418
||||| ||||||| ||||||| ||||||| ||||||| |||||||
794 T...GGCAGATATACATATATATTC 816
||||| ||||||| ||||||| ||||||| ||||||| |||||||

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305 GluMetVal.....HisProValAlaGluArgLeuAsp.....11 316
      |||.....|||.....|||
6324 CAAGTTTTCATGACACACATATGACAGACAGTACACAGATGACAAAG 6373
      |||.....|||.....|||
316 eleMetSerLeuValLeuSerTyrMetLysAspValCysTyrValAspG 333
      |||.....|||.....|||
6374 CTATATCAACATATATGTTAACCCACACGACAAAGTTG.....C 6411
      |||.....|||.....|||
333 LysValAspAsnGlyLysThrLysAspLeuTyrArgAspLeuLea 349
      |||.....|||.....|||
6412 CGCAGCTATGATGCTGCTTACGCTCTTATCTTCTTACCAATATATGTA 6461
      |||.....|||.....|||
350 LLePheAspLysLeuLeu.....LeuProth 358
      |||.....|||.....|||
6462 GTTAAACACAAACATATTCACATTCGCTCTTACTGTACATATTTACA 6511
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358 rHisAlaSerCysHisValGln..... 365
      |||.....|||.....|||
6512 GTACCGAATATGATATGAACTTACGATCTGCTGACGCTGTCAG 6561
      |||.....|||.....|||
365 ..... 365
6562 ATAAAGACTCAATTTCTTCTCTCTGTCAGGCTTATTCCTGCTTG 6611
      |||.....|||.....|||
366 PhePheMetPheTyrLeuCysSerPheLys.....LeuGlyPheAlaG 380
      |||.....|||.....|||
6612 TATTACACATATATCTCTGATGATGACCAATTCCTTACGTTGTTGA 6661
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380 uAlaPheLeuGluHisLeuTyrLysLysLeuGlnAspProSerAsnPro 397
      |||.....|||.....|||
6662 GCGAGAGGTAAAGATTATTTGTCGAAAGTACGAGATTCTCAAGTATA 6711
      |||.....|||.....|||
397 lAlaLeuValGlnAlaAlaCysAspTyrLecGlySerPheLeuAlaArg 413
      |||.....|||.....|||
6712 GCTTCCTCAACACAAATGATATGATATCTCAATTCCTTTTAGGGAAA 6761
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414 AlAlaPheTyrLeuProLeuLeuValLysSerCysLeuAspLeuLeu 430
      |||.....|||.....|||
6762 TTGACAGAGCT.....TTAGGTTTACGCTGATTTGATTAAT 6799
      |||.....|||.....|||
430 lAsnTyrLeuHisLLeuTyrLeuAsnAsnGlnAspSerGlyThrLysAla 447
      |||.....|||.....|||
6800 CGACTTAAACAT..... 6812
      |||.....|||.....|||
447 heCysAspValAlaLeuHisLysProPheTyrSerAlaCysGlnAlaVal 463
      |||.....|||.....|||
6812 ..... 6812
464 PheTyrThrPheValPheArgHisLysGlnLeuSerGlyAspLeuLeu 480
      |||.....|||.....|||
6813 .....CAGATTCAATCA.....GTCAA 6829
      |||.....|||.....|||
480 sGluAlaTyrLeuGlnTyrLeuGlnSerLeu.....AsnPheGluArgLLe 494
      |||.....|||.....|||
6830 GGAAGGCTTATTTGCTTAACATCATTCATTCATTTTCACAAAGCT 6879
      |||.....|||.....|||
495 .....ValMetSerGlnLeuAsnProLeu 502
      |||.....|||.....|||
6880 ATGATGACCATGATGAACCTTCTGCTTATATGAAAGAGTTTCGTGATG 6929
      |||.....|||.....|||
503 LysLLeCysLeuProSerValValAsnPhePheAlaAlaLeuThrAsnL 519
      |||.....|||.....|||
6930 GCATACAAAGCTGAGTATGATTCATTCATTCGCTGCTATCTCATCC 6979
      |||.....|||.....|||
519 s.....TyrGlnLeuValPheCysTyrThrLLeLecGluArgAsnAsn 534
      |||.....|||.....|||
6980 ACTCTGATACAAAGCTCTTGGATTTCTGAAGTCTTGAG...AAATATA 7026
      |||.....|||.....|||
534 rGlnMetLeuProValLLeArgSerThrAlaGlyLysAspSerValGln 550
      |||.....|||.....|||
7027 AGCTTGTAAATAAGTGTGTGTCGAGACATTCGAAGACCAACATTTGAA 7076
      |||.....|||.....|||
551 lLeCysThrAsnProLeuAspThrPhePheProPheAspProCysValLe 567
      |||.....|||.....|||

```

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7077 GTTACTGTGATGAA..... 7094
      |||.....|||.....|||
567 uLysArgSerLysLysPheLLeAspProTyrGlyTyrGlnValTyrGluAsn 584
      |||.....|||.....|||
7094 TSCAAAGACACAACTTATATGATGACCAATTTTTCAGCTTATATCTAAA 7144
      |||.....|||.....|||
584 eleSerAlaGluGluLeuGlnGluPheLysLysProMetLysLysAspLLe 600
      |||.....|||.....|||
7144 GACCAACACAGAAATGACAGGTTTTCAGATATACATA..... 7181
      |||.....|||.....|||
601 ValGluAspLLeAspAspAspPheLeuLysGlyGluValProGlnAsnAs 617
      |||.....|||.....|||
7182 .....GATGAATTAAGACATAAATCTTCTGAGATCAATGATGATCTTAA 7225
      |||.....|||.....|||
617 P.....ThrValLLeGlyLLe..... 622
      |||.....|||.....|||
7226 TGTCAATTCAAATGTTGGCATGGCAATATGAGCAATATACACTAGTAA 7276
      |||.....|||.....|||
623 .....TheProSerPheAspThrHis 630
      |||.....|||.....|||
7276 AAGAGATTATCAATATCCAGAGTACCTCTGCTTTCATGATGAT 7322
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seq_name: /cqn2-6/ptodata/1/ina/5R.COMB.seq:US-08-680-627-1
seq_documentation_block:
? Sequence 1, Application US/08680327
? Patent No. 5859421
? GENERAL INFORMATION:
? APPLICANT: Staskiewicz, Brian S., Oldroyd, Giles Edward,
? APPLICANT: Salmeron, John M., Romeros, Carlos
? TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR PLANT
? TITLE OF INVENTION: PATHOGEN RESISTANCE
? NUMBER OF SEQUENCES: 15
? CORRESPONDENCE ADDRESS:
? ADDRESSER: Klarnoust Sparkman Campbell Lotah &
? STREET: One World Trade Center
? STREET: 121 S.W. Salmon Street
? STREET: Suite 1600
? CITY: Portland
? STATE: Oregon
? COUNTRY: United States of America
? ZIP: 97204
? COMPUTER READABLE FORM:
? MEDIUM TYPE: Disk, 3-1/2 inch
? COMPUTER: IBM PC compatible
? OPERATING SYSTEM: MS DOS
? SOFTWARE: Wordperfect 5.1
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/08/680,327
? FILING DATE: July 11, 1996
? CLASSIFICATION: 800
? PRIOR APPLICATION DATA:
? APPLICATION NUMBER: 08/410,912
? FILING DATE: September 22, 1994
? CLASSIFICATION: 800
? APPLICATION NUMBER: 08/227,360
? FILING DATE: April 13, 1994
? ATTORNEY/AGENT INFORMATION:
? NAME: Dow, Alao, E.
? REGISTRATION NUMBER: 35,124
? REFERENCE/DOCKET NUMBER: 5151-45078
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: (503) 226-7391
? TELEFAX: (503) 226-9446
? INFORMATION FOR SEQ ID NO: 1:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 5475 base pairs
? TYPE: nucleic acid
? STRANDEDNESS: double stranded
? TOPOLOGY: linear

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WATTE

THE ALGEBRAIC THEORY OF LOGIC

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alignment_scores:      101.00      length:      69%
                        quality:      0.357      gaps:      2
Percent Similarity:    40.486      Percent Identity: 16.45%

alignment_block:
US-09-932-678-2 x US-09-228-246-3 ..

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Ratio:	0.387	Gaps:	25
Percent Similarity:	45.455	Percent Identity:	19.755

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alignment_block:
US-09-942-678-2 x US-09-080-897-3
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Align seq 1/1 to: US-09-080-897-3 from: 1 to: 4378

1 MetIlaIaIaProLeuLeuHisThrArgLeuProGly.....AspAlaI 15
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15 AlaSerSerSeraIaValLysLysLeuGlyAlaSerArgThrGlyIles 32
486 GAGCTCTAGCTCTGGCATGATGTACATCCAGCAGCTGAGGTGGGGCTGC 535
32 eraSnmel.....ArgAlaLeuGluAsnAspIhe 42
536 GGGATATCGACCTGTGACCTGCTTGAATCCCTTCGACGCTCTCCAC 585
43 AsnSerProProArgLysThrValArgPheGly.....GlyTh 55
586 AATAACCTGTGCATTTGGGTGGCAGACATTGGTGGCTGAGAGCCTAGCTC 635
55 rValThnGluValLeu.....LeuLysTrpLysGlyGluThr.... 68
636 CTATTGGACATCTCCAAACGCTCATGATGAGAAAGAGAGACACTTCTG 685
69AsnAspIheGluLeuLeuLysAsnGluLeu 79
686 GAACTACGACAGCGCGAAACCGACATGAGATGATATCGCC.....TGTGG 729
80 AspProAspIleLysAspAspGlnIleLeuSerTrpLeuLeuGluPheArg 96
730 AAGCGTTTCATGACACACACACATTGGATCCAAACATATGTTGGAGACGGA 779
96 gSerSerIleMetTrpLeuThrLys..... 104
780 AGAAGAGATCTACTGCTGGTGCACAGCCATGATGATCTGCTGTCGCCATA 829
105AspPheGluGlnLeuIleSerIleLeuAlaGluProTrp 118
830 TGATGATGATCGACCAACAAAGCTGCTCTGCTCCCTCTGATCTGCGCCAG 879
119LeuAsnArg.....Ser 122
880 CCGGAGGACATGATGATGAACGAGATTTCTAGAGCGCAATGACACAGAGCTCA 929
122 rGlnThrValaGluGluTrpLeuAlaPheLeuGluGlyAsnLeuValSera 139
930 AATGATGATGAGCTGCAACGCTTCCAGGCACCTTGTGAGCGATTAAAAAGTG 979
139 lagGlnThrValaPheLeuArg.....ProGlyLeuSerMetIleAlaSerHis 154
980 GAACTCTATTGCGCTCAAGTGGCATGCTCTAAGCTCATCAATGCTCTC 1029
155 PheValProProArg..... 159
1030 ATCACTCGAGCTGAGGAACTGTGATTCCGAGAGCTTCGAGAGATGAA 1129
159ValIleIleLysGluGlyAspVal 167
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168 AspValSerAspSerAspAspGluAspAsnLeuProAlaAsnIleHis 184
1180 GATTCTCTGATCTGACAGGAGCGGCTGATGATATGCTGCTGGAGATGGA 1229
184 pThrCysHisArgAlaLeuGlnIleIleAlaIarGlyValProSerThr. 200

1 230 TCACTTGGTCAGCTTTTCAATATATTTTAAACACGCTCAACATTTCA 1279
201 ProPhePheMetProLeuValGluHisPheProHis 214
1280 AGGCACAGCCACACCTCCGCTCT AATCGACACCTCTCTCTG 1428
215 ValArgLysSerGluArgThrLeu G1 223
1324 GTCGCAAAATGATTAAGAACCGACGACACGACATCAATAAACATGATTCAGA 1478
223 TCYSTYValHisAsnLeuLeuArgHisLeuSerValTyrPheProThrLeu 240
1374 AGCTGTTCTCGAAATAGTCTTACACAAAATGGAAGCTATCTTACTTCA 1423
240 TGGHSGluLeuLeuLeuLeuLeuLeuLeuLysLeuLysAspVal 256
1424 AATGCCGACACCTCGACATGATATTCACACTG GTTCATCA 1467
257 AspAlaSerArgGluGlyLeuLysAspAlaGluThrAlaThrGlu 273
1468 ATGATTGATTAACCAACAGTGGAAAAATCTGAGGCCAAAGCTACAGACT 1517
27 TYSGLYGLYThrAspSer ThrGlu 282
1518 GCAAAAAAAGCTGGATTCAGTAATTAACACACCGGCACAGTTTAACCTAG 1567
242 TyleuPheAsnMetAspGluAsp GluThrGluHisGluThrLys 297
1567 AATGCAAAAACATGCAAAATGACTTTGAGCACAACACTTCATATCTTCA 1617
256 AlaGlyProGluArgAlaGluAspMetValHisProValAlaGluArgLe 314
1618 GAGAAAGAGATAGGTTGATTTGATTTGAAAAAGAGATCACTCAACAGAA 1667
314 uasplleuMetSerLeuValLeuSerTyrMetLysAspValLysTyr 341
1666 ACAAGACCTCGACGGCAGAGCTTCCACAGCTACACGAAAGCTTCCCAAG 1717
331 alaAspLysValAspAsnGlyLysTyrLysAspLeuTyrArgAspLe 347
1718 TTTCAAAAACACTACAGAAATGCAAGAAATGAAATGGCTTCCTCTCTCT 1767
347 u LeuAsnLeuPheAspLysLeuLeuLeuProThrHisA 360
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366 LAsertyshLysAlaGlnPhe PheMetPheTyrLeuGlySerPheLys 375
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376 LeuGlyPheAlaGluAlaPheLeuGluHisLeuTyrPheLysLeuHis 392
1866 CTCTCTCTCTCTGAGAGTGTCTGCTCCACT 1897
392 ProSerAsnProAlaLeuLeuArgGluAlaAlaLysAsnTyrIleLys 409
1897 1897
409 CT 425
1898 CCCCCTCTCT 1909
426 LeuAspLeuValAsnTyrPheLeuHisLeuTyrLeuAsnGluHisAsp 442
1910 CTTCAGCTACTGTATCTC CTCCACTCTCTCTCT 1942
442 TlyThrLysAlaPheGlyAspValAlaLeuHisGlyProPheTyrSerA 459
1943 TACTGAGAGTGGTGTATAC CCCCCTGCTCTCA 1976
456 LAGCTGAlaVal 463
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257 AsnAlaSerArgGlnGlyIleGluAspAlaGluIuThrAlaThrGlnTh 274
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1468 ATGATTGATAAACAAGGTGGAAATCTGAGCCAAAGCTACAGAGCT 1517
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273 cysglycylThrAspSer.....ThrGlnG 282
      :          :          :          :          :          :
1518 GCAAAAAAGATTCATTCAGATTAACAGCGCGGACAGCTACAGCTAG 1567
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282 IyLeuPheAsnMetAspGluAsp...GluGluThrGlnHisGluThrLys 297
      :          :          :          :          :          :
1568 AATGCAAAAAAGATGAAATGAGCTTTAGAGCAAACTTCAGGATCTTCA 1617
      :          :          :          :          :          :
298 AlaGlyProGlnArgLeuAspGlnMetValHisProValAlaGluArgLe 314
      :          :          :          :          :          :
1618 GGAATAAAGCATGCTGCTGATCTTGAATAAGCAGACATCAGCTGACATCA 1667
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314 uAspIleLeuMetSerLeuValLeuSerTyrMetLysAspValcysTyrV 331
      :          :          :          :          :          :
1668 ACAAGACCTTGACGCGACAGGTGTCAGAGCTGACAGAGAGCTTCCAGG 1717
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331 aIAspGlyLysValAspAsnGlyLysThrLysAspLeuTyrAlaGlyAsp 347
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1718 TGTCAAAAGACATGAGATGCCAGATGCCAAGATGAGCTTCTCTCTGCT 1767
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347 u.....IleAsnIlePheAspLysLeuLeuProThrHisA 360
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1768 GTGTTGTTGACCTCTGTTTCTAGCAGAGTGTCTCTTCCCTTCCCTCC 1817
      :          :          :          :          :          :
360 LaSerCysHisValGlnPhe...PheMetPheTyrLeuCysSerPheLys 375
      :          :          :          :          :          :
1818 TCCTTCGCGCTGTCAGCTGTCGACCTTATGTCAGCTCCCGCCAGCCGAC 1867
      :          :          :          :          :          :
376 LeuGlyPheAlaGlnAlaPheLeuGlnHisLeuTyrLysLysLeuGln 392
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1868 CTCTCTCTCTGAGGTGTGTCTCCACCAT..... 1897
      :          :          :          :          :          :
392 ProSerAspProAlaIleLeuArgGlnAlaAlaGlyAsnTyrIleGly 409
      :          :          :          :          :          :
1897 ..... 1897
      :          :          :          :          :          :
409 erPheLeuAlaArgAlaLysPheIleProLeuIleThrValLysSerCys 425
      :          :          :          :          :          :
1898 .....CCCCCTCT.....TGC 1909
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426 LeuAspLeuLeuValAsnTyrPheHisIleLeuLysAsnGlnAspSe 442
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1910 CTCGAGTACTTATATCC...CTCCACCTCTCTCT..... 1942
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442 rGlyThrLysAlaPheCysAspValAlaLeuHisGlyProPheTyrSer 459
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1944 .....TACCTGAGAGTCTCTGTATAC...CCCCCTCCCGCCA 1976
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459 IacysGlnAlaVal 463
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1977 GTTCCTGCTGCTG 1990

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seq.name: /cqn2_6/piodata/1/lna/6A-COMB.seq:05-08-899-595-2

seq.documentation_block:

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: Sequence 2, Application US/0889595
: Patent No. 6111072
:
: GENERAL INFORMATION:
: APPLICANT: Natsumiya, Shuh
: APPLICANT: Takahashi, No. 6111072uaki
: TITLE OF INVENTION: RHO TARGET PROTEIN HUMAN MDIA AND GENE
: NUMBER OF SEQUENCES: 14
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Foley & Lardner
: STREET: 3000 K Street, N.W., Suite 500
: CITY: Washington
: STATE: D.C.
: COUNTRY: USA

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ZIP: 20007-5199
:
: C-MEDIA: REPRODUCIBLE FORM:
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: MEDIUM TYPE: Floppy disk
:
: COMPUTER: IBM PC compatible
:
: OPERATING SYSTEM: PC-DOS/MS-DOS
:
: SOFTWARE: Patent Release #1.0, Version #1.40
:
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/899,595
: FILING DATE: 24-JUL-1997
: CLASSIFICATION: 435
: PRIORITY APPLICATION DATA:
: APPLICATION NUMBER: JP 8-242701
: FILING DATE: 26-AUG-1996
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: JP 9-90170
: FILING DATE: 25-MAR-1997
: ATTORNEY/AGENT INFORMATION:
: NAME: Stephen A. Bent
: REGISTRATION NUMBER: 29,768
: REFERENCE/WORK NUMBER: 049441/0112
:
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (202)672-5300
: TELEFAX: (202)672-5399
:
: INFORMATION FOR SEQ ID NO: 2:
:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 4909 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: Molecule type: cDNA to mRNA
:
: ORIGINAL SOURCE:
: ORGANISM: Mouse
:
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 94..3858
:
: US-08-899-595-2

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Quality	100.50	Length	572
Ratio	0.387	Gaps	25
Percent Similarity	45.455	Percent Identity	19.755

alignment_block:

US-09-932-678-2 x US-08-899-595-2 ..

Align seq 1/1 to: US-08-899-595-2 from: 1 to: 4899

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15 aAlaSerSerAlaValAlaLysLysLeuGlnGlyAlaSerArgThrGlyLys 42
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486 GAGCTCTACGCTCTGATATGATGATATGATGATGATGATGATGATGATG 545
      :          :          :          :          :          :
32 erAsnMet.....ArgAlaLeuGlnuAsnAspPhe 42
      :          :          :          :          :          :
536 GCAATATGCAAGCTGTTAGTGTGCTTGAGTCCCTTGAGCTGCTGTAAC 585
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586 AATAACCTTCAATTTGGTGCAACATTTGGTGCTGAGAGCTTAGGCTAG 645
      :          :          :          :          :          :
55 rValThrGlnValLeu.....LeuLysTyrLysGlyGluThr.... 68
      :          :          :          :          :          :
636 CTATTGAGCAATCTTAACAGACTTCATATGATAAAGAGAGAGACTTCTG 685
      :          :          :          :          :          :
69 .....AsnAspPheGlnLeuLeuLysAsnGlnLeuLeu 79
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686 GAACTAGCAAGCTGAAACCAACATGATGATATGCTCC.....TCTTTG 729

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80 AspGluAspIleLysAspAspIleIleIleAsnThrPheLeuLeuPheAsn 96
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700 AAGCTTCTTCAGAAATGAAATTTGGTAATCAAAATATGTTGGAGACAGCA 779
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96 GAGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 104
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
780 AGAAGAGAAATCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 829
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105 ..... AspGluGluIleGluLeuLeuLeuLeuLeuLeuLeuLeuLeu 118
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840 TGAATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 879
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
119 ..... LeuAsnAsnG..... 122
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
880 CTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 929
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122 TGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 139
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940 AATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 979
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149 TATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 154
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980 GATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1029
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159 TbeValProGluG..... 159
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
1030 ATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1079
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
159 GATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 159
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1080 GATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1129
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166 ..... ValIleIleGlySerGluGlyAspVal 167
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1130 ATTAACATATGAAATGAAATGATGATGATGATGATGATGATGATGATGATG 1179
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168 AspValIleAspGluAspAspGluAspAspGluProAlaAspGluAsn 184
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184 TbeGlySerGluGluGluGluGluGluGluGluGluGluGluGluGluGluGlu 200
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
1230 TGAATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1279
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
201 ..... ProGluPheGluMetProIleGluValIleGlyPheProGlu 214
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1280 AGGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1323
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
215 ValAcGlySerGluGluGluGluGluGluGluGluGluGluGluGluGluGlu 223
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
1324 GTCGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1373
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225 GTCGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 240
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1474 ATGCTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1423
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240 TGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 256
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1424 ATGCTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1467
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257 AsnAlaSerAlaGluGluGluGluGluGluGluGluGluGluGluGluGluGlu 273
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
1498 ATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1517
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273 TGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 282
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282 TGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 297
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1568 AATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1617
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298 AlaGlyProIleGluGluAspIleMetValIleSerValAlaGluGluGlu 314

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1618 GATAAATAAATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1667
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414 GATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 431
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1668 AATAAATAAATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1717
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
431 AlAspGlyValAspAsnGlyLysPheLysAspLeuGlyArgAspGlu 447
   |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
1718 TGTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1767
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447 G..... IleAsnIlePheAspIleGluGluGluProIleHis 460
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1768 GTCATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1817
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409 GTPheGluAlaGluAlaGluAlaGlyPheIleProGluIlePheValLysSerGly 425
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1918 GTCGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 1942
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442 TGTGlyThrLysAlaPheGlyAspValAlaGluGluHisIleGlyProGluGlySerGlu 459
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459 TAspGluAlaVal 463
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seq_name: /gen2/6/pdata/alpha/1/ind/68.00MB.seq:US-08-974.462.2
seq_documentation_block:
Sequence 2: Application US/08974462B
Patient No. 6191270
GENERAL INFORMATION:
APPLICANT: DOBILHE, PIERRE
APPLICANT: DADBERSIES, PIERRE
TITLE OF INVENTION: MALARIAL PRE-ERYTHROCYTIC STAGE POLYPEPTIDE MOLECULES
FILE REFERENCE: 0660-0125-0 PCT
CURRENT APPLICATION NUMBER: US/08/974.462B
CURRENT FILING DATE: 1998-02-06
EARLIER APPLICATION NUMBER: PCT/FR94/00894
EARLIER FILING DATE: 1994-06-12
EARLIER APPLICATION NUMBER: FR 95/07007
EARLIER FILING DATE: 1995-06-14
NUMBER OF SEQ ID NOS: 29
SUBJECT: Patient In Vet. 2.0
SEQ ID NO: 2
LENGTH: 5361
TYPI: ONA
ORIGIN: P. falciparum
FEA: ORG:
NAME/KEY: CDS
LOCUS: IN: (1) ..(5461)
US OR: 54462.2
alignment_statistics:
Quality: 99.50
Length: 559

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Ratio: 0.381 Gaps: 23
Percent Similarity: 46.691 Percent Identity: 19.499
alignment block:
US-09-932-678-2 x US-08-973-462-2 ..
Align seq 1/1 to: US-08-973-462-2 from: 1 to: 5361

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seq_documentation_block:
? Sequence 4, Application US/08776265
? Patent No. 6001631
? GENERAL INFORMATION:
? APPLICANT: BLANCHET, Francis
? APPLICANT: CAMERON, Beatrice
? APPLICANT: CROUZET, Joel
? APPLICANT: FAMECHON, Alain
? APPLICANT: FERRERO, Lucia
? TITLE OF INVENTION: No. 6001631el Topoisomerase IV, Corresponding
? NUMBER OF SEQUENCES: 17
? CORRESPONDENCE ADDRESS:
? ADDRESSEE: Finnegan, Henderson, Parabow, Garrett &
? ADDRESSEE: Dunnot, L.L.P.
? STREET: 1300 I. Street, N.W., Suite 700
? CITY: Washington
? STATE: D.C.
? COUNTRY: USA
? ZIP: 20005-3315
? COMPUTER READABLE FORM:
? MEDIUM TYPE: Floppy disk
? OPERATING SYSTEM: IBM PC compatible
? SOFTWARE: Patent In Release #1.0, Version #1.30
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/08/776,265
? FILING DATE: 24-JAN-1997
? CLASSIFICATION: 435
? ATTORNEY/AGENT INFORMATION:
? NAME: Elnaudi, Carol P.
? REGISTRATION NUMBER: 32,220
? REFERENCE/DOCKET NUMBER: 03806,0394-00000
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: (202) 408-4000
? TELEFAX: (202) 408-4444
? INFORMATION FOR SEQ ID NO: 4:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 2402 base pairs
? TYPE: nucleic acid
? STRANDEDNESS: single
? TOPOLOGY: linear
? MOLECULE TYPE: cDNA
? US-08-776-265-4

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Ratio: 0.462      gaps: 19
Percent Similarity: 45.319      Percent Identity: 17.660
Alignment_block:
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Align seq 1/1 to: US-08-776-265-4 from: 1 to: 2402

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seq_documentation_block:

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? Sequence 13, Application US/08417340A
? Patent No. 5719057
? GENERAL INFORMATION:
? APPLICANT: HADJINAM, KAREN
? APPLICANT: LE BOUQUELLES, HEATRICE
? APPLICANT: WHITTING, PAUL
? TITLE OF INVENTION: STABLY TRANSFECTED CELL LINE EXPRESSION
? TITLE OF INVENTION: GABA-A RECEPTOR AND NOVEL CLONED
? NUMBER OF SEQUENCES: 20
? CORRESPONDENCE ADDRESS:
? ADDRESSEE: J. MARK HAND - MERCK & CO., INC.
? STREET: 126 EAST LINCOLN AVENUE - P.O. BOX 2000
? CITY: RAHWAY
? STATE: NJ
? COUNTRY: US
? ZIP: 07065-0900
? COMPUTER READABLE FORM:
? MEDIUM TYPE: Diskette
? COMPUTER: IBM Compatible
? OPERATING SYSTEM: DOS
? SOFTWARE: FASTESTO for Windows Version 2.0
? CURRENT APPLICATION DATA:
? APPLICATION NUMBER: US/08/417,330A
? FILING DATE: 05-APR-1995
? CLASSIFICATION: 435
? PRIOR APPLICATION DATA:
? APPLICATION NUMBER:
? FILING DATE:
? ATTORNEY/AGENT INFORMATION:
? NAME: HAND, MARK
? REGISTRATION NUMBER: 36,545
? REFERENCE/DOCKET NUMBER: 111091A
? TELECOMMUNICATION INFORMATION:
? TELEPHONE: 908-594-3905
? TELEFAX: 908-594-4720
? TELEX:
? INFORMATION FOR SEQ ID NO: 13:
? SEQUENCE CHARACTERISTICS:
? LENGTH: 1638 base pairs
? TYPE: nucleic acid
? STRANDEDNESS: double
? TOPOLOGY: linear
? MOLECULE TYPE: cDNA
? PEZIORE:
? NAME/KEY: Coding Sequence
? LOCATION: 87...1562
? OTHER INFORMATION:
? US-08 417-340A-13

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alignment_scores:

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Quality: 97.00 Length: 626
Ratio: 0.406 Gaps: 31
Percent Similarity: 38.179 Percent Identity: 18.051

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alignment_block:

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US-09-932-678-2 x US 08-417-340A 13
Align seq 1/1 to: US-08 417-340A 13 from: 1 to: 1638

```

```

124 AsnLeuValSerAlaGlnThrValIlePheLeuArgProCysLeuSerPheI 151
      |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:

```


[illegible]

• • •

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 30, 2002, 02:33:51 : Search time 2612.22 Seconds
(without alignments)
16566.781 Million cell updates/sec

Title: US-09-932-678-1
Perfect score: 2068
Sequence: 1 acaagagctgtgctggaag.....tgacatttgattcccaat 2068

Scoring table:
IDENTITY_NUC
Gapop 10.0, Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues
Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :
1: gb_da:*
2: gb_hg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_pro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vi:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_om:*
21: em_of:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pl:*
26: em_ro:*
27: em_sts:*
28: em_un:*
29: em_vi:*
30: em_hg_hum:*
31: em_hg_inv:*
32: em_hg_other:*
33: em_hgq_inv:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
------------	-------	-------------	--------	-------	-------------

1	2068	100.0	2068	9	AF227156	AF227156 Homo sapi
2	2020	97.7	3756	9	HS2272050	AX046047 Sephiro
3	2018.4	97.6	2040	6	AX036047	AX012547 Homo sapi
4	1211.6	58.6	135873	2	AC012547	AF014444 Homo sapi
5	1211.6	58.6	192815	9	AC013444	AK055742 Homo sapi
6	567	27.4	2493	9	AC055742	BC009198 Homo sapi
7	506.4	24.5	1177	6	AC009198	AX274407 Sephiro
8	420.8	20.3	1461	6	AX274407	BC009441 Homo sapi
9	420.8	20.3	2627	9	BC006441	AC040158 Homo sapi
10	208	10.1	129473	2	AC040158	AC092137 Homo sapi
11	208	10.1	143900	2	AC092137	AC007615 Homo sapi
12	192	9.3	175691	2	AC007615	AC009130 Homo sapi
13	192	9.3	191496	2	AC009130	AC0092562 Homo sapi
14	192	9.3	208008	2	AC0092562	AC092475 Homo sapi
15	190.8	9.2	187643	2	AC092475	AC017077 Homo sapi
16	188.8	9.1	173166	9	AC092475	H0AF001549 Homo chr
17	188.8	9.1	179150	2	AC017077	AC106788 Homo sapi
18	188.8	9.1	202004	9	H0AF001549	AC092137 Homo sapi
19	187.2	9.1	170611	9	AC106788	AC092137 Homo sapi
20	158	7.6	143900	2	AC009130	AC092475 Homo sapi
21	153.2	7.4	191496	2	AC009130	AC092475 Homo sapi
22	153.2	7.4	195413	2	AC092475	AC092475 Homo sapi
23	153.2	7.4	247331	2	AC092475	AC092475 Homo sapi
24	146.4	7.1	2112	3	AY061123	AY061123 Homo sapi
25	140.6	6.8	179150	2	AC017077	AY061123 Homo sapi
26	131	6.3	293	9	HS15183F	AY061123 Homo sapi
27	127	6.1	78347	2	AC022642	AY061123 Homo sapi
28	125.4	6.1	53814	2	AC107769	AY061123 Homo sapi
29	118.6	5.7	53814	2	AC107769	AY061123 Homo sapi
30	115	5.6	824	9	HS2272050	AY061123 Homo sapi
31	108.6	5.3	821	9	HS2272050	AY061123 Homo sapi
32	91.4	4.4	108040	2	AC068150	AY061123 Homo sapi
33	91.4	4.4	171930	2	AC092137	AY061123 Homo sapi
34	91.4	4.4	173166	9	AC092137	AY061123 Homo sapi
35	91.4	4.4	316296	2	AC092137	AY061123 Homo sapi
36	70.8	3.4	78347	2	AC022642	AY061123 Homo sapi
37	62.8	3.0	405	11	G27379	AY061123 Homo sapi
38	62.8	3.0	6287	2	AC017809	AY061123 Homo sapi
39	62.8	3.0	96018	3	AE004783	AY061123 Homo sapi
40	62.8	3.0	172748	3	AC018484	AY061123 Homo sapi
41	62.8	3.0	176424	3	AC011757	AY061123 Homo sapi
42	59.6	2.9	174531	2	AC023752	AY061123 Homo sapi
43	51.4	2.5	125020	9	AF429415	AY061123 Homo sapi
44	48.2	2.3	8797	6	AF027419	AY061123 Homo sapi
45	48.2	2.3	10338	4	BC004586	AY061123 Homo sapi

ALIGNMENTS

RESULT 1
LOCUS AF227156 Homo sapiens RRNA, complete cds.
DEFINITION AF227156
ACCESSION AF227156.1 GI:7670099
KEYWORDS
SOURCE
ORGANISM human.
human.
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

REFERENCE
1 (bases 1 to 2068)
Moorefield, B., Greene, E. A. and Reeder, R. H.
RNA polymerase I transcription factor Rn3 is functionally
conserved between yeast and human
Proc. Natl. Acad. Sci. U.S.A. 97 (9): 4724-4729 (2000)
2 (bases 1 to 2068)
Moorefield, B., Greene, E. A. and Reeder, R. H.
Direct Submission
Submitted (20-JAN-2000) Basic Sciences, Fred Hutchinson Cancer
Research Center, 1100 Fairview Avenue N., Seattle, WA 98199, USA

JOURNAL MEDLINE
REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES

Db	1481	CAGTATTAGAGTACGGCTGGAGGAGACTAGTCGACATCTGCACAAACCCCGCTGACA	1740
QY	1741	ccctctccctcttgatccctgtgtgtgtgaagagtcgaagaatctcatctctatc	1800
Db	1741	ccctctctccctcttgatccctgtgtgtgtgaagagtcgaagaatctcatctctatc	1800
QY	1801	atcaagatggaagaagatgaatgctgaagaagctacagaagatctgaagaaccaga	1860
Db	1801	ATCAGGTTGGGAAGACATGATGCTGGAAGAGCTACAGAGCTTCAGAAATCCATGAAA	1860
QY	1861	aggaaatgatagaagatgaagatgtagactctctctgaaggcgaaatgcccgaatgata	1920
Db	1861	AGGACATGATGGAAATGATGAATATATGACTTTCTGAAGGCGAATGCGCCAGATGATA	1920
QY	1921	ccgtgatatggaatcacacccaagctctcttgacacgcattccgaagctcttaagatg	1980
Db	1921	CCGTGATTGGGATGACACCAAGCTCTCTTGCACGCAATTCGGAAGTCTTCAAGTAGTG	1980
QY	1981	tggatcccccccgctatgtacatgaagaccagtcggctcttgacagagaataattgaga	2040
Db	1981	TGGGTTCCCAACCCGCTGTGTGACATGCAACCCAGTCCGCTCTGACGCGAGAAATTTGGA	2040
QY	2041	ctgaatgataacatttggaattccccat	2068
Db	2041	CTGACATGTGACATTTGSGATTCCGCAT	2068
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LOCUS	HS4272050		
DEFINITION	HS4272050	3756 bp	mRNA linear
ACCESSION	AJ272050		
VERSION	AJ272050.1	GI:10046713	
KEYWORDS	TIF-1A; transcription initiation factor.		
SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Eukariota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
AUTHORS	1 (bases 1 to 3756) Bodem,J., Hoffmann-Rohrer,U., Ross,W., Delius,H., Vingron,M. and Grunert,I.		
TITLE	Cloning and functional characterization of transcription initiation factor TIF-1A, a growth-dependent regulator of ribosomal RNA synthesis		
JOURNAL	unpublished		
REFERENCE	2 (bases 1 to 3756)		
AUTHORS	Bodem,J.J.		
TITLE	Direct Submission		
JOURNAL	Submitted (07-FEB-2000) Bodem J.J., Molekulare Biologie der Zelle		
FEATURES	11. Deutsches Krebsforschungszentrum, INF.280, 69120 Heidelberg, GERMANY		
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	23..3756		
	/gene="TIF-1A"		
	23..1978		
	/gene="TIF-1A"		
	/function="transcription initiation factor"		
	/evidence="experimental"		
	/product="transcription initiation factor 1A protein"		
	/protein_id="CAC07955.1"		
	/db_xref="GI:10046714"		
	/translation="MAAFLLHTRLPGDAASSAVKKIGASRTGISNNRALEDNFNS		
	PKPTVRGGTIVTEVLLKRRKGINDDELKNQLDDPKDDITNNLIEPRSSIMYI		
	TKDPELIVSLRLPMLNRSQTVVEEYLAIGNIVSAQTVLRGLSMASHVYPRKS		
	1IKKDVIVSDSDDEDNDLPANFDICHAALDILIRYVSTWFLMLILVEFPVPRKS		
	KRTLEGVNHLIRLIRISVYKPTLRHEILELLEKLLKIDYNASRQIGDAEFAATGCG		
	TDSTEDLNNMDEFTFEHEIKAGGEEDQWYPAERLDTLMSLVLSMKQVCTVDGG		
	VDNKTCDLVNDLNIIDKLLLPTHASQVDFMFYLCSPFVGLGFAEALFELMKRLDQ		

[illegible]

[illegible][illegible]

Db 97266 GATTATCTTCTGACATTTTGACACGCGATACACCTTACAAATTAACAGCATATAT 97325
 QY 657 qfaccatgacacgagcgglllclcalqccaaiaacgagggaaatltccattgllcga 716
 Db 97326 GTCCGACTGGCAGCTGGTGTCTTTTGGCCATACCTGTGGAAACATTTCCATTGTGCT-- 97383
 QY 717 aaatcgagagacacatgaatgaatllacglttaacttaactgaagatgaatgaatltt 776
 Db 97384 -AATCGAGAGAGACACTGGAATTTTACATTTACATTTACTTAAGATTAATATACCTTT 97442
 QY 777 ccaactllagacatgaatcttgaagcttattatgaagaactacacatllcgaatllcga 836
 Db 97443 CCAACCTTGAAGAGATGAATTTTGAAGCTTATTATTAAAAATCTATTCAATTTGATGTC 97502
 QY 837 aatgcatcccgagagagatllgaagatgcltgaagaacagcaactcaactllatgag 896
 Db 97503 AATATCATTTGCGCATGATTTTGAGATGCTGAGAGAAACAGCAGCTTAACCTTGTGCTG 97562
 QY 897 aacgagtcacagagagatglttattatgaagatgaagatgaagaactgaactgaaga 956
 Db 97563 ACACATTCATGGAATGATTTTCAATATGACGAGATGAACAGC-----TGTTCCT 97616
 QY 957 aagagcggllcctlaaacgagcgcacacagatgaagcctgaagcgcgcgcgcgcgcac 1016
 Db 97617 AAGCTGACACTGCAATGGCTGGTGCATGCTGACCTGACCTGACCTGACCTGACCTGAC 97676
 QY 1017 ctgaatgcltllgalltllcttcaatgaagagatgcltctatg-----tagatllgaag 1071
 Db 97677 CTGCTGCTCTTCTGTTTGAGAGTACTGGAATGATGCTGCTATGTAATAATACATGTAAG 97736
 QY 1072 ttatatacggaacaaagagatctatctgcgactgaataacatctcttgaagaactcc 1131
 Db 97737 TTGATACACAAACAAATTAAGATTTATA-----ATATATCACAACCTGATTAACATA 97788
 QY 1132 tcttgcacacacacatgctctcgcacatgaaglllclcalqccaaiaacgagggaaatltcc 1191
 Db 97789 TTGACAAACACCTGCTGCTGCTGACGATGAGTTTATGTTTATGCTGCTGCTGCTGCT 97848
 QY 1192 aatllgagatllcgaagagcaatlllgaagaactcctgq-----aaaaatllcgaagac 1248
 Db 97849 AATTGGACACTGCAAGAGCTTTTGGACCTCTGCGGAAAAAATAATGACAGATCCAA 97908
 QY 1249 gtaatcctgcacatcagcagcagcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgc 1308
 Db 97909 ATATATCTGTCATCATAGGTGAGCTGCGGAATTAATATGGAACCTCTTTGGCAAAAG 97968
 QY 1309 claaatlalctctcttatt---actgaaatcagcgcgcgcgcgcgcgcgcgcgcgcgcgcgc 1365
 Db 97969 CTAAATTGATCTCTCTCATTTACAACTATAAATCATGCCAGTCTTTGCGTTAACTGCC 98028
 QY 1366 tgcacatatacctlaataaccagatllcgaagaacagcaatllcgaagatllcgcctcc 1425
 Db 98029 CACACAAATACCTTACTTAACAGAGATCTGAGATGAAGGCTTTTGATATATGCTGCTGC 98088
 QY 1426 atggaccattllactcagc 1485
 Db 98089 ATCAAAATTTTACTGAGCTTCTCAAGCTGTCTGCCACACCTTTGTTTATGACACAAAG 98148
 QY 1486 agctlltgaagcgaacacclgaagaagatllcgaagatllcgaagatllcgaagatllcga 1545
 Db 98149 AACTTTTAACGCTGAACCTGAAGAGTTTGAAGATCTCTGACGCTGCTGATATTTTAAG 98208
 QY 1546 agatlltgaagcgaacacclgaagaagatllcgaagatllcgaagatllcgaagatllcga 1605
 Db 98209 GGAATAGTACGAGCTGAAGCTGAATATTTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 98268
 QY 1606 ttgctgaac 1665
 Db 98269 TTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 98325
 QY 1666 atggc 1725
 Db 98326 AATAGACTGAG 98368

QY 1726 caaacccgctgagacacctcttcccccttgaatctctgagatgaagatgaagatgaagat 1785
 Db 98369 CCAACCAATGAGAGCTCTTCCCTTTGATCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 98428
 QY 1786 tcatgacatctatctatctatctatctatctatctatctatctatctatctatctatctat 1845
 Db 98429 TCATTGATTTTATTATTCATATGCAAGATGCAAGATGCAAGATGCAAGATGCAAGATGCAAGAT 98488
 QY 1846 agaaacccac 1905
 Db 98489 ACAAACCTATGAAAG 98545
 QY 1906 tgc 1960
 Db 98546 TTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 98605
 QY 1961 ccaagctcttgaagatgaagatgaagatgaagatgaagatgaagatgaagatgaagatga 2019
 Db 98606 CCAAAATGCTTGAAGTACTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 98665
 QY 2020 tctgaagcgaagatlltgaagatgaagatgaagatgaagatgaagatgaagatgaagatga 2075
 Db 98666 TCTCATCTGAGAGATTTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 98701

RESULT 5
 AC013444
 LOCUS Homo sapiens BAC clone RP11-470B22 from 2, complete sequence.
 DEFINITION AC013444
 ACCESSION AC013444.8 GI:18093267
 VERSION AC013444.8
 KEYWORDS HTG.
 SOURCE human.
 ORGANISM Homo sapiens.
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1
 AUTHOR(S) Sulston, J.E. and Waterston, R.
 TITLE Toward a complete human genome sequence.
 JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
 MEDLINE 99063792

REFERENCE 2
 AUTHOR(S) Wallington, S.K., Baekenson, W. and Spalding, L.
 TITLE The sequence of Homo sapiens BAC clone RP11-470B22
 JOURNAL unpublished (2001)
 REFERENCE 3
 AUTHOR(S) Waterston, R.H.
 TITLE Direct Submission
 JOURNAL Submitted (11-NOV-1999) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA
 4 (bases 1 to 192815)

REFERENCE 4
 AUTHOR(S) Waterston, R.
 TITLE Direct Submission
 JOURNAL Submitted (09-JAN-2002) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
 On Jan 9, 2002 this sequence version replaced gi:13518264.
 COMMENT -- Genome Center

Center: Washington University Genome Sequencing Center
 Center code: WUGSC
 Web site: <http://genome.wustl.edu/asg>
 Contact: sapiens@wustl.wustl.edu
 Summary Statistics
 Center project name: H_NH0470B22

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:

all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality > 40); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/usc>

SOURCE INFORMATION:

The RP111 human PAC library was made from the blood of one male donor as described by Osoegawa, K., Moon, P. Y., Zhao, B., Penangu, E., Hattori, M., Cateforis, J., and de Jong, P. J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 54:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Victor de Jong (<http://bartsch.med.tufts.edu>)

VERTICAL PHRED 6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is AC024616; the clone sequenced to the right is RP1119023. Actual start of this clone is at base position 1 of RP111470822; actual end is at base position 192815 of RP111470822.

Unread over homopolymeric run from base position 164727 to 164753.

FEATURES:

Source: location/coordinates

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/db_xref "taxon:9606"

/chromosome "2"

/map "2"

/clone "RP111470822"

/clone_lib "RP111"

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611..878

/rpt_family "ERV1"

879..1032

/rpt_family "(L1A)n"

1046..1092

/rpt_family "ERV1"

1987..2446

/rpt_family "L2"

2737..2972

/rpt_family "L1"

3702..4749

/rpt_family "AT-rich"

4870..4891

/note "match to EST BC071256 (NID:q841643)"

6880..6992

/rpt_family "MIR"

8677..8934

/rpt_family "Alu"

8773..8800

/rpt_family "AT-rich"

8955..8969

/rpt_family "AT-rich"

9207..9247

/rpt_family "L2"

9442..9773

/rpt_family "L2"

9977..10011

/rpt_family "(L1A)n"

10976..11265

/rpt_family "Alu"

12178..12209

/rpt_family "(T)n"

12362..12390

/rpt_family "AT-rich"

12702..12741

/rpt_family "(16)n"

12731..12758

/rpt_family "(GA)n"

12758..12785

/rpt_family "(T)n"

13033..13054

/rpt_family "(AAAA)n"

13195..13699

/rpt_family "L1"

13543..13667

/rpt_family "AT-rich"

14142..14305

/rpt_family "MIR"

14500..14534

/rpt_family "AT-rich"

14617..14735

/rpt_family "MIR"

14736..14836

/rpt_family "MER2-1ypc"

14837..15106

/rpt_family "Alu"

15107..15317

/rpt_family "MER2-1ypc"

15318..15345

/rpt_family "MIR"

15578..16066

/rpt_family "ERV1"

16072..16112

/rpt_family "AT-rich"

18192..18222

/rpt_family "AT-rich"

18612..18899

/rpt_family "Alu"

19181..19485

/note "match to EST BC0991 (NID:q95918) yq02104.s1"

19879..21714

/note "similar to Rattus norvegicus EST Y16973 (NID:q4727035)"

19890..20284

/note "match to EST BC09197 (NID:q954024) yq02104.c1"

19915..20369

/note "similar to Homo sapiens EST BC35986 (NID:q13627431)"

19923..20378

/note "similar to Homo sapiens EST AA213789 (NID:q1812416)

20311..21

19923..20369

/note "similar to Homo sapiens EST BC428405 (NID:q1334811)"

20091..20378

/note "similar to Homo sapiens EST A1742153 (NID:q5110441)

20158..20664

/note "similar to Homo sapiens EST AV703279 (NID:q10720608)"

20243..21062

/note "similar to Mus musculus EST BC295610 (NID:q13057417)"

20412..21044

/note "similar to Homo sapiens EST BC734554 (NID:q10148346)"

20567..20666

/note "similar to Homo sapiens EST BC778654 (NID:q14048951)"

20592..20963

/note "similar to Homo sapiens EST AA481295 (NID:q2210847)

20703..21362

/note "similar to Homo sapiens EST BC502871 (NID:q1464388)"

20798..21056


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FEATURES
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    /db_xref="taxon:9606"
    /chromosome="16"
    /clone_lib="Caltech human BAC library A"
    /clone_id="31241 g 35543 t 1468 others"

BASE COUNT      30715 a 10506 c 31241 g 35543 t 1468 others
ORIGIN

Query Match
Best Local Similarity 10.1%, Score 208; DB 2: Length 129473;
Matches 208; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1861 agagatagttgaaatgaatgaatgaatttttaaaagagaaatgcacccaatgata 1920
Db 123967 AGGAGATAGTGTGAAGATGTAAGATGATGATTTCTTAAGAGCCAGATGGCCAGATGATA 124046

QY 1921 cgtatattggagatcaccaaatcctttgacagcatttcgaatttcctaaatgag 1980
Db 124047 CCGTATTGGAGATCCACATCCAACTCTTTGACACCGCATTTCCGAATGCTTTAAAGTAGTG 124106

QY 1981 ttagctcccccacccgattgttcatcatgcaacccatccctcttgaagcagaatatttqta 2040
Db 124107 TGGCTCCGCCACCCGTTGTTGACATGCAACCCAGATCCCTCTGACGCGCAATTTGTGA 124166

QY 2041 ctgaatattgacatttggattccccat 2068
Db 124167 CTGAGATGTGACATTTGGATTTCCCAT 124194

RESULT 11
LOCUS AC092137/c 143900 bp DNA linear HTG 22-JUN-2001
DEFINITION Homo sapiens chromosome 16 clone RP11-344H15, WORKING DRAFT
AC092137
VERSION AC092137.1 GI:14522983
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
  1 (bases 1 to 143900)
  DOE Joint Genome Institute.
  Sequencing of Human Chromosome 16
  Unpublished
  2 (bases 1 to 143900)
  DOE Joint Genome Institute.
  Direct Submission
  Submitted (22-JUN-2001) Production Sequencing Facility, DOE Joint
  Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
  -----Genome Center
  Center: Joint Genome Institute
  Center Code: JGI
  Web site: http://www.jgi.doe.gov
  -----
  Project Information
  Center Project Name: 538644
  Center clone name: RPC1-11_344H15
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  Summary Statistics
  Consensus quality: 124071 bases at least Q40

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Consensus quality:	146140 bases at least Q20
Estimated insert size: <td>211000; adaptor-tp estimation</td>	211000; adaptor-tp estimation
Estimated insert size: <td>142200; sum-of-coverage estimation</td>	142200; sum-of-coverage estimation
Quality coverage: <td>4.0 in Q20 bases; adaptor-tp estimation</td>	4.0 in Q20 bases; adaptor-tp estimation
Quality coverage: <td>5.94 in Q20 bases; sum-of-coverage estimation</td>	5.94 in Q20 bases; sum-of-coverage estimation
* NOTE: This is a 'working draft' sequence. It currently	
* consists of 18 contigs. The true order of the pieces	
* is not known and their order in this sequence record is	
* arbitrary. Gaps between the contigs are represented as	
* runs of N, but the exact sizes of the gaps are unknown.	
* This record will be updated with the finished sequence	
* as soon as it is available and the accession number will	
* be preserved.	
1	1034; contig of 1033 bp in length
1034	1134; gap of unknown length
1134	2547; contig of 1414 bp in length
2548	2647; gap of unknown length
2648	4205; contig of 1558 bp in length
4206	4305; gap of unknown length
4306	6564; contig of 2259 bp in length
6565	6664; gap of unknown length
6665	9065; contig of 2401 bp in length
9066	9165; gap of unknown length
9166	12059; contig of 2894 bp in length
12060	12159; gap of unknown length
12160	19295; contig of 7136 bp in length
19296	19395; gap of unknown length
19396	25688; contig of 6273 bp in length
25689	25768; gap of unknown length
25769	31013; contig of 5245 bp in length
31014	31113; gap of unknown length
31114	36555; contig of 5442 bp in length
36556	36655; gap of unknown length
36656	45338; contig of 8883 bp in length
45339	45438; gap of unknown length
45439	52918; contig of 7480 bp in length
52919	53018; gap of unknown length
53019	65720; contig of 12702 bp in length
65721	65820; gap of unknown length
65821	76109; contig of 10289 bp in length
76110	76209; gap of unknown length
76210	85439; contig of 9240 bp in length
85440	85539; gap of unknown length
85540	101620; contig of 16081 bp in length
101621	101720; gap of unknown length
101721	122163; contig of 20443 bp in length
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Matches 208; Conservative 0; Mismatches 0; Gaps 0;	
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27279	CGGATATGGAATGACACCAAGTCTCTTGAACGATCTTCGAAGTCTTCGATATG 27219
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* 144469 144569: gap of unknown length
* 144569 163180: contig of 18612 bp in length
* 163181 163280: gap of unknown length
* 163281 176573: contig of 13293 bp in length
* 176574 176674: gap of unknown length
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* 185463 185562: gap of unknown length
* 185563 187643: contig of 2081 bp in length.

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Best Local Similarity 94.38; Pred. No. 3.6e-41;
Matches 198; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

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DB 40557 ACACCGATATAGTGCACAGATGAGATGATGACTTCTGAAAGCTCAAGTGCCTACATGA 40616
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VERSION AJ272050.1 GI:10046713
KEYWORDS TIF-1A; transcription initiation factor.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
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Bodem J., Hoffmann-Kohrer U., Koss W., Delius H., Vingron M. and
Grunmt J.
TITLE Cloning and functional characterization of transcription initiation
factor TIF-1A, a growth-dependent regulator of ribosomal RNA
synthesis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 3756)
AUTHORS Bodem J.J.
TITLE Direct Submission
JOURNAL Submitted (07-FEB-2000) Bodem J.J., Molekulare Biologie der zelle
II, Deutsches Krebsforschungszentrum, INF.280, 69120 Heidelberg,
GERMANY
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[illegible]

[illegible][illegible]

[illegible]

Genome, Mark Keitelman and Anuradha Madan

Clone distribution: M6E clone distribution information can be found through the F.M.A.G.E. Consortium/URL at: <http://image.fnl.gov>

Series: 1841, Release: 24 Row: K Column: 14
This clone was selected for full length sequencing because it passed the following selected criteria: matched mRNA g1: 3355302.

Location/Qualifiers

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/clone_1b "NH-M6E-8"

/lab host "DH10B-K"

/note "Vector: pCR17"

316, 546

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ORIGIN

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Ratio: 4.714 Gaps: 2

Percent Similarity: 74.242 Percent Identity: 74.485

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586 TTTTTCATTTTTCATTTTTCATTTTTCATTTTTCATTT 645

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646 TTTTTCATTTTTCATTTTTCATTTTTCATTTTTCATTT 665

157 TTTTTCATTTTTCATTTTTCATTTTTCATTTTTCATTT 174

686 TTTTTCATTTTTCATTTTTCATTTTTCATTTTTCATTT 745

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seq name: 3d, 10:AY061123

seq document location block:

DEFINITION AY061123

VERSION AY061123.1

KEYWORDS

ORIGIN

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Sequence submitted by:

Lawrence Berkeley National Laboratory

Lawrence Berkeley National Laboratory

Lawrence Berkeley National Laboratory

Lawrence Berkeley National Laboratory

Lawrence Berkeley National Laboratory

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FEATURES

Location/Qualifiers

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178. 2013
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BASE COUNT 649 a 405 c 414 g 644 t
ORIGIN

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Ratio: 1.967 Gaps: 18
Percent Similarity: 60.395 Percent Identity: 30.501

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35 GAlaLeuGlUasnaSPhePheasnsErProArGlyThrValArp 52
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337 CGAAATTTTCATCTGTCGAAGAGATTCACATCTCTCCGGGAACGACA 386
82 pIleIyAspAspGlnIleIleasnTrpIleuLeuGluPheArGSerI 99
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437 TTCATATATTGACTCCAGAAATTTGTAACCTGTGCGAGCGCTTACTGTC 486
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587 CTAAACTAATTTGCTACATGATCTC.....GGA 615
166 AspValAspValSerAsp.....SerAspAspGluAspAspAsnLeu 180
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180 oAlaAspPheAspThrCysHisArqAlaLeuGluIleIleAlaArqTyr 197
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247 lGleGluLysLeuLeuLysIleuAspValAsnAlaSerArqGlnGlyIle 263
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305 GluMetValHisProValAlaGluArqLeuAspIleLeuMetSerLeuG 321
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336 sEAsnGlyLysThrLysAsp...LeuTyrArqAspLeuIleasnIlePhe 351
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352 AspLysLeuLeuLeuProThrHisAlaSerCysHisValGlnPheMet 368
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368 tPheTyrIleuLysSerPheLysLeuGlyPheAlaGluAlaPheLeuGlu 385
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[illegible]

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seq_description: block:			
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Accession	271927		
VERSION	271927.1 GI:1404567		
KEYWORDS	RRM3 gene.		
FEATURES	RRM3 yeast.		
ORIGIN	Source: Flycos. Cerevisiae		
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ORIGIN:6001..12000	Source: Flycos. Cerevisiae		
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seq_documentation_block:

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 ACCESSION AC007615
 VERSION AC007615.6 GI:14589428
 KEYWORDS HTG; HTGS_PHASE2; HTGS_DRAFT; HTGS_ACTIVEFIN.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo;
 1 (bases 1 to 175691)
 DOE Joint Genome Institute.
 JOURNAL Sequencing of Human Chromosome 16
 REFERENCE Unpublished
 2 (bases 1 to 175691)
 Bruce,D., Mundt,M., Doggett,N., Munk,C., Saunders,E., Robinson,D.,
 Jones,M., Buckingham,J., Chasteen,L., Thompson,S., Goodwin,L.,
 Bryant,J., Tesmer,J., Meincke,L., Longmire,J., White,S., Tatum,O.,
 Campbell,C., Fawcett,J., Meltbie,M., Bussod,M., Sutherland,R.,
 McMurry,K., Han,C. and Deaven,L.
 DIRECT SUBMISSION
 Submitted (20-MAY-1999) Center for Human Genome Studies, DOE Joint
 Genome Institute, Los Alamos National Laboratory, MS M888, Los
 Alamos, NM 87545, USA
 COMMENT On Jul 3, 2001 this sequence version replaced gi:13928651.

Sequence Quality Assessment:
 This entry has been annotated with sequence quality
 estimates computed by the Phrap assembly program.
 All manually edited bases have been reduced to quality zero.
 Quality levels above 40 are expected to have less than
 1 error in 10,000 bp.
 Base-by-base quality values are not generally visible from the
 Genbank flat file format but are available as part
 of this entry's ASN.1 file.

Sequence Quality Assessment:

This entry has been annotated with sequence quality
 estimates computed by the Phrap assembly program.
 All manually edited bases have been reduced to quality zero.
 Quality levels above 40 are expected to have less than
 1 error in 10,000 bp.
 Base-by-base quality values are not generally visible from the
 Genbank flat file format but are available as part
 of this entry's ASN.1 file.

Genbank flat file format but are available as part
 of this entry's ASN.1 file.

* NOTE: This is a 'working draft' sequence. It currently
 * consists of 1 contigs. Gaps between the contigs
 * are represented as runs of N. The order of the pieces
 * is believed to be correct as given, however the sizes
 * of the gaps between them are based on estimates that have
 * provided by the submitter.

* This sequence will be replaced
 * by the finished sequence as soon as it is available and
 * the accession number will be preserved.
 * 1 175691: contig of 175691 bp in length.
 * location/qualifiers

FEATURES

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BASE COUNT 42392 a 42125 c 45277 g 45896 t 1 others
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 393 osGAspProAlaIleIlePArgGlnAlaGlyAsnTyrIleGlySerP 410
 30055 AAGTATTCCTGGCATCATCAGGAGAGCTGTGTGAAATTAATATGGAGCT 30104
 410 heLeuAlaArgAlaIlePheLeuLeu..... 420
 30105 TTTTGGCAAGCAGTAATTAATTCCTTAATGTAAGTACCTAATTTTCT 30154
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 30405 GTGTTGATATAGTTTGTATGTTTGTAGATTTTGTCAATTT 30454
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 30455 GCAATCCAAATGATATGTAATGTATTAAGCTGAGAAAGCCCTTTAAA 30504

XX Particularity: Tumors

XX Claim 1: Fig. 2: 48pp: German.

XX This invention describes a novel DNA sequence (1) that encodes the RNA polymerase I transcription factor TIF-1A which has anti-tumor,

anti-proliferative and proliferative inducing activity. The invention also

describes (1) DNA (1a) encoding a protein (11) with the biological

activity of TIF-1A; (2) a ribozyme (8) corresponding to (1) or (1a) and

able to bind specifically to, and cleave, its transcribed RNA so as to

reduce or inhibit synthesis of the corresponding protein; (3) an

antisense RNA (AS) with binding properties similar to R; (4) an

expression vector that contains (1), (1a) or sequences that encode R or

AS; (5) host cells containing the vectors of (4); (6) TIF-1A or (11)

encoded by (1) or (1a); (7) preparation of TIF-1A or (11) by culturing

cells of (6); (8) ligands that bind to TIF-1A or (11); (9) a diagnostic

method for detecting abnormal TIF-1A expression; and (11) kit for

carrying out the method in (10); (1), and similar sequences that encode

proteins with equivalent activity, expression vectors containing them, or

their expression products are used to treat or prevent disorders

associated with reduced cellular proliferation, to stimulate cellular

proliferation, and to promote tissue regeneration, e.g., after injury or

radiation therapy. Ribozymes, antisense sequences directed against (1),

also ligands and antagonists of TIF-1A, are used to treat or prevent

disorders associated with excessive cellular proliferation and to inhibit

proliferation, especially in treatment of cancers. (1) and specific

ligands for TIF-1A (particularly antibodies (Ab)) are also useful for

diagnosis of altered TIF-1A expression by (in)direct determination of the

concentration, length and/or sequence of TIF-1A or its mRNA, e.g., for

detecting mutations. Ab can also be used for immunoprecipitation of

TIF-1A and for isolation of related sequences from cDNA expression

libraries. (1) allows recombinant production of TIF-1A in sufficient

quantities for therapeutic use.

XX Sequence: 2040 bp; 563 A; 444 C; 474 G; 568 T; 2 other;

XX

XX

XX

XX

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XX

XX

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XX

XX

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XX

XX

XX

XX

XX

XX

XX

XX

[illegible][illegible]

RESULTS

(1) AA'92255 standard; (10NA; 2110 BP

XX	AA552255
AC	

XX
DT 14-PTA-2002 (first entry)

XX [1N], encoding novel human c

XX Haplotype chromosome mapping: gene mapping: gene location
KM

XX food supplement; medical imaging; diagnostic; genetic disorder; etc.

105 HETERO SAPIENS.
XX

PN	W0200175007-A2
XX	

FD 11-887 2001
XX

40-MAR 2001; 2001W0-US08631.
 PF
 XX

PR 41-MAR-2000; 2000US-0540217
PR 24-AUG-2000; 2000US-0649167

XX
1A (HYSF-) HYSFQ INC.

XX
P1
DROGAS RI, LUG, TANG YU

XX
10K
W11: 2000-6-8 852/73.

DR P-PSI/B; ABC'28068,
XX

New isolated poly
diisotics, fore

responsible
biodiversity

Classification: SEO ID No 28059; 10400; English.

The invention relates to isolated polynucleotide (1) and

polypeptide (11) sequences. (11) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromo-

of gene mapping, and in recombinant production of (11). The nucleotides are also used in diagnostics as expressed sequence tags

(c) for identifying expressed genes, (I) is useful in gene therapy techniques for identifying expressed genes, (II) or to treat disease states involving loss of normal activity of (III) or to treat disease states involving

quantitative polymerase chain reaction (PCR) as molecular weight markers and as

equivalent to a 100% increase in the amount of food supplement. (11) and its binding partners are useful in medical treatment of sites expressing (11) and (11) are useful for treating

disorders involving aberrant protein expression or biological activity. The polynucleotide sequences have applications in

diagnostics, forensics, gene mapping, identification of mutations

CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPo
 CC at http://wipo.int/pub/publ/published_pcl_sequences.

XX
 CC Sequence 2410 BP; 672 A; 437 C; 547 G; 754 T; 0 other;

Query Match 10.9%; Score 639.4; DB 23; Length 2410;
 Best Local Similarity 96.8%; Pred. No. 1.4e-177;
 Matches 706; Conservative 0; Mismatches 16; Indels 7; Gaps 5;

QY 1347 gatcttttgaatacactgacacatacattacaaacagatctcgaacaaagca 1406
 DB 1 gatcttttgaatacactgacacatacattacaaacagatctcgaacaaagca 60
 QY 1407 ttctgagatgtgtctccatgacacatttaccacgtcgcgaagatgtttacacc 1466
 DB 61 ttctgagatgtgtctccatgacacatttaccacgtcgcgaagatgtttacacc 120
 QY 1467 ttatgtttgaacacagacgtctttgaacgcaaacctgaagaagtttgaacatctt 1526
 DB 121 ttatgttttgaacacagacgtctttgaacgcaaacctgaagaagtttgaacatctt 180
 QY 1527 caaagctgaatttgaacgataatgatacgcacgaataacccctgaagat-ttacct 1585
 DB 181 caaagctgaatttgaacgataatgatacgcacgaataacccctgaagatgtacct 240
 QY 1586 gacctgagttgtaactttttgtgtcaatcacaataatgatacagctgtcttctgcta 1645
 DB 241 gacctgagttgtaactttttgtgtcaatcacaataatgatacagctgtcttctgcta 300
 QY 1646 caacatattgaagagacacatcgcacagatgctgcagatcttttgaagctgagag 1705
 DB 301 caacatattgaagagacacatcgcacagatgctgcagatcttttgaagctgagag 360
 QY 1706 agactcagatgacatctgacacaaacccgagacacttctcccttt-gatcctctga 1764
 DB 361 agactcagatgacatctgacacaaacccgagacacttctccctttgatacctctgag 420
 QY 1765 tgcctgaagaggtcaagaatc-cattgactctatc---atcagatctgaagaacatg 1820
 DB 421 tgcctgaagaggtcaagaatc-cattgactctatc---atcagatctgaagaacatg 480
 QY 1821 agtctgaagaggtcagagatctcaagaacacccatgaagaagagagatgagaa 1880
 DB 481 agtctgaagaggtcagagatctcaagaacacccatgaagaagagagatgagaa 540
 QY 1881 gataatgactttcgaagaagcgaagtgccccaagaaalalacagatgattgatacaca 1940
 DB 541 gataatgactttcgaagaagcgaagtgccccaagaaalalacagatgattgatacaca 600
 QY 1941 agctcctttgaacagatcttccgaagatccttcaagttagttgagctccccaacccatgtt 2000
 DB 601 agctcctttgaacagatcttccgaagatccttcaagttagttgagctccccaacccatgtt 660
 QY 2001 taag-ttgaacccagctccctctgaagcacaataattgtgatactatgaagattggag 2059
 DB 661 taag-ttgaacccagctccctctgaagcagaataattgtgatactatgaagattggag 720
 QY 2060 attcccccatt 2068
 DB 721 attcccccatt 729

RESULT 6
 AAS94253
 ID AAS92253 standard; CDNA; 776 BP.
 XX
 AC AAS92253;

XX
 DT 13-FEB-2002 (first entry)
 XX
 DE DNA encoding novel human diagnostic protein #28057.
 XX
 KW Human: chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostics; genetic disorder; ss.
 XX
 OS Homo sapiens.
 XX
 PN W0200175067-A2.
 XX
 PD 11-JUL-2001.
 XX
 XX 30-MAR-2001; 2001WO-0508631.
 XX 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX
 PA (HVSF-) HVSFO INC.
 PT Dmanac RT, Liu C, Tang YF.
 XX
 DR MP1: 2001-639362/73.
 DR P-FSD8: AHC28056.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 XX
 XX Claim 1; SEQ ID NO 28057; 103pp; English.

XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging or sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biochemical activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPo
 CC at http://wipo.int/pub/publ/published_pcl_sequences.

XX
 CC Sequence 776 BP; 206 A; 157 C; 198 G; 215 T; 0 other;

Query Match 27.5%; Score 568.6; DB 23; Length 776;
 Best Local Similarity 97.8%; Pred. No. 5.6e-157;
 Matches 629; Conservative 0; Mismatches 9; Indels 5; Gaps 5;

QY 4 gaagcttgaatgaagaagatgaagatccagctgaagcgaagcagatcagattatctg 63
 DB 133 gaagcttgaatgaagaagatgaagatccagctgaagcgaagcagatcagattatctg 192
 QY 64 gacccaatgagacacagctgcttccacag-catttgcgaggaatgagagccttctctc 122
 DB 133 gacccaatgagacacagctgcttccacagcttcttgcgaggaatgagagccttctctc 252
 QY 124 tctgcaattgaagacagcagctctc-gaagactgaagattccaatatatgatacattaga 181
 DB 253 tctgcaattgaagacagcagctctcagctcctgaagactgaagattccaatatatgatacattaga 412

07	182	gaatgaatcttttcaattctctctcccaaaacacacgatttcgatttgcgtgaacatgcacaca	284
08	415	gaatgaatcttttcaattctctctcccaaaacacacgatttcgatttgcgtgaacatgcacaca	372
09	242	aattcttctgaattacacaaagaggttgaaacaaatgaatttgatttattgaagacacagct	301
10	478	aattcttctgaattacacaaagaggttgaaacaaatgaatttgatttattgaagacacagct	432
07	402	gtttgaattcccaattcaaaaggttgaacacacacacacacacacacacacacacacacac	359
10	448	gtttgaattcccaattcaaaaggttgaacacacacacacacacacacacacacacacacac	492
09	600	tctatctatctatcttgacaaagagcttttgagacacattatcattatattaaagatgcct	418
10	463	tctatctatctatcttgacaaagagcttttgagacacattatcattatattaaagatgcct	552
09	419	ttaatttgaattacaaattcaaacacacacacacacacacacacacacacacacacacac	478
10	558	ttaatttgaattacaaattcaaacacacacacacacacacacacacacacacacacacac	612
09	613	atctac	598
10	613	atctac	672
09	613	atctac	598
10	613	atctac	732
09	613	atctac	641
10	748	tattcttcttgcacatttctgaacacacacacacacacacacacacacacacacacacac	775
RESULTS			
7			
AN	AN587643	AN587643 standard; cDNA: 1260 bp.	
AN	AN587643		
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
RNA encoding novel human diagnostic protein #23447.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	
10	155	FEBS 2002 (first entry)	
Human: chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss.			
07	155	FEBS 2002 (first entry)	
08	155	FEBS 2002 (first entry)	
09	155	FEBS 2002 (first entry)	

[illegible]

Db 793 atgaaccatcatgatactatcttctctactatcttgaatgtaacttctcaagaataatgta 852
 Qy 715 gaaatcatgagagaacacacgcggaaatgtaactgtaacttaactgaagtaatgataat 774
 Db 853 atcaatgaataaataatgaatgtaataatgaataaagctgagcttctacatgatat 912
 Uy 775 tt 776
 Db 913 tt 914

RESULT 8
 ID AAS80955/c
 AAS80955 standard; cDNA: 655 bp.
 AC
 XX AAS80955:
 XX
 DT 13-FEB-2002 (first entry)
 XX
 DE DNA encoding novel human diagnostic protein #16750.
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 OS Homo sapiens.
 XX
 PN W0200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PF 30-MAR-2001; 2001MO-US08631.
 XX
 PR 41-MAR-2000; 2000MS-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 PR
 PA (HYSEQ.) HYSEQ INC.
 XX
 PI Dymaeac RT, Liu C, Tang YT;
 XX
 DR WP1: 2001-639362/73.
 DR P-PSDB: ABG16768.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 XX
 PS Claim 1; SEQ ID No 16759; 103pp: English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at http://wipo.int/pub/published_pat_sequences.
 XX
 XX Sequence 655 BP: 207 A; 134 C; 115 G; 199 T; 0 other;

[illegible]


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RESULT 10
ABAR3038
ID ABAR3038 standard; DNA: 1461 BP.
AC ABA83038;
XX
DE 05-FEB-2002 (first entry)
XX
DEF Human transcription factor TRFX-65 coding sequence.
XX
KW Human; transcription factor; TRFX; cell proliferative disease;
KM autoimmune disease; inflammation; neurological disease;
KM developmental disorder; cancer; AIDS; infection; cytostatic; anti-HIV;
KM neuroprotective; antinflammatory; gene therapy; ds.
OS Homo sapiens.
PN WQ20017277-A2.
PD 04-OCT-2001.
PE 13-MAR-2001; 2001WQ-US08117.
PR 13-MAR-2000; 2000US-0188986.
PX
PY (JINCY-) INCYTE GENOMICS INC.
PA Hillman JL, Baughn MR, Yue H, Lal P, Lu DM, Patterson C;
PI Azimzal Y, Bandman O, Tang YT, Mathur P, Shah P, Au-Yang J;
P1 Reddy R;
XX
DR WP1: 2001-570896/64.
DRI P-PDB: ABR50214.
PT Novel transcription factor polypeptides, used to treat diseases
PI associated with altered activity and expression of TRFX, and to screen
PP for agents capable of modulating its activity -
PS Claim II; Page 299; 327pp; English.
XX
CC The present sequence is the coding sequence for a human transcription
CC factor. The transcription factor and its coding sequence are useful in
CC the diagnosis, treatment and prevention of diseases associated with
CC altered expression of the transcription factor e.g. cell proliferative,
CC autoimmune/inflammatory, neurological and developmental disorders. A
CC number of specific disorders/diseases are given in the specification,
CC including: arteriosclerosis, cirrhosis, hepatitis, cancers, AIDS,
CC allergies, anaemia, asthma, autoimmune thyroiditis, bronchitis, atopic
CC dermatitis, diabetes mellitus, emphysema, Goodpasture's syndrome, gout,
CC Grave's disease, multiple sclerosis, osteoarthritis, pancreatitis,
CC psoriasis, rheumatoid arthritis, systemic lupus erythematosus, ulcerative
CC colitis, uveitis, Alzheimer's disease, Huntington's disease, Parkinson's
CC disease, stroke, and viral, bacterial, fungal and protozoal infections.
XX
SQ Sequence 1461 BP; 413 A; 335 G; 298 C; 415 T; 0 other;
```

QY	1468	1469	1470	1471	1472	1473	1474	1475	1476	1477	1478	1479	1480	1481	1482	1483	1484	1485	1486	1487	1488	1489	1490	1491	1492	1493	1494	1495	1496	1497	1498	1499	1500
QY	1468	1469	1470	1471	1472	1473	1474	1475	1476	1477	1478	1479	1480	1481	1482	1483	1484	1485	1486	1487	1488	1489	1490	1491	1492	1493	1494	1495	1496	1497	1498	1499	1500
Db	710	711	712	713	714	715	716	717	718	719	720	721	722	723	724	725	726	727	728	729	730	731	732	733	734	735	736	737	738	739	740	741	742
QY	1428	1429	1430	1431	1432	1433	1434	1435	1436	1437	1438	1439	1440	1441	1442	1443	1444	1445	1446	1447	1448	1449	1450	1451	1452	1453	1454	1455	1456	1457	1458	1459	1460
Db	770	771	772	773	774	775	776	777	778	779	780	781	782	783	784	785	786	787	788	789	790	791	792	793	794	795	796	797	798	799	800	801	802
QY	1468	1469	1470	1471	1472	1473	1474	1475	1476	1477	1478	1479	1480	1481	1482	1483	1484	1485	1486	1487	1488	1489	1490	1491	1492	1493	1494	1495	1496	1497	1498	1499	1500
Db	850	851	852	853	854	855	856	857	858	859	860	861	862	863	864	865	866	867	868	869	870	871	872	873	874	875	876	877	878	879	880	881	882
QY	1518	1519	1520	1521	1522	1523	1524	1525	1526	1527	1528	1529	1530	1531	1532	1533	1534	1535	1536	1537	1538	1539	1540	1541	1542	1543	1544	1545	1546	1547	1548	1549	1550
Db	890	891	892	893	894	895	896	897	898	899	900	901	902	903	904	905	906	907	908	909	910	911	912	913	914	915	916	917	918	919	920	921	922
QY	1568	1569	1570	1571	1572	1573	1574	1575	1576	1577	1578	1579	1580	1581	1582	1583	1584	1585	1586	1587	1588	1589	1590	1591	1592	1593	1594	1595	1596	1597	1598	1599	1600
Db	950	951	952	953	954	955	956	957	958	959	960	961	962	963	964	965	966	967	968	969	970	971	972	973	974	975	976	977	978	979	980	981	982

RESULT 11
 ID AAX51643
 AC AAX51643 standard; cDNA: 437 bp.
 XX
 AC AAX51643:
 XX
 DT 21-JUN-1999 (first entry)
 XX
 DE Human secreted protein 5' EST Sbj ID No:242.
 XX
 KW Human secreted protein; EST; expressed sequence tag; diagnosis;
 KW genetic; gene therapy; chromosome mapping; signal peptide;
 KW intram regulatory sequence; cytokine activity; cell proliferation;
 KW differentiation; hematopoiesis; regulation; tissue growth regulation;
 KW retroactive hormone regulation; chemokine; chemokine; haemostatic;
 KW thromolytic; anti-inflammatory; tumour inhibition; ds.
 XX
 OS Homo sapiens.
 XX
 PN W06905549-A2.
 XX
 PD 11-FEB-1999.
 XX
 PE 31-JUL-1998; 98WO-1801231.
 XX
 PR 01-AUG-1997; 97US-0905279.
 XX
 PA (GIST) GENSET.
 XX
 PA DeJfert A, Dumas Mline Edwards J, Lacroix B;
 PL
 DR WPI: 1999-153739/13.
 DR P-ESDR: AAY128A5.
 XX
 PT New nucleic acids encoding human secreted proteins obtained from
 PT cDNA libraries derived from testis, ovary, uterus and spleen tissue
 XX
 (Claim 1: Page 344-345; 562pp; English).
 XX
 CC AAX51459 to AAX51631 represent 5' expressed sequence tags (ESTs) for
 CC human secreted proteins, and encode the proteins given in AAY12641 to
 CC AAY12713, respectively. The proteins given represent the signal peptide
 CC and an N-terminal fragment of a secreted protein. The nucleic acid
 CC sequences can be used for producing secreted human gene products. They
 CC can also be used to develop products for diagnosis and therapy. The
 CC proteins obtained may have cytokine activity, haematopoiesis regulating
 CC proliferation/differentiation activity, haematopoiesis regulating
 CC activity, tissue growth regulating activity, reproductive hormone
 CC regulating activity, chemokine/chemokine activity, haemostatic and
 CC thromolytic activity, receptor/ligand activity, anti-inflammatory
 CC activity, tumour inhibition activity or other activities. The products
 CC can be used in diagnostic, gene therapy and chromosome mapping procedures

10

the sequences can also be used for obtaining corresponding promoter sequences. The nucleic acids encoding the signal peptide can be used for direct or indirect selection of a polypeptide or the insertion of a polypeptide into a membrane, or importing a polypeptide into a cell.

50. Sequence 447 BP; 118 A; B⁵ C²; 102 G; 131 W; 1 other;

Quincy March	29, 48; Street 420; 108 20; 1004th 437;
--------------	---

Matchless 4.24; Conservative 1; Mistmatch 6;

by the effect of a single q value. The value of q is 172

CUIC

1. *Introduction*

2033 quacccacqct qtt baad ccacqacat caacqat cat caac qccct qct. 352

110 182 Graduate Catalogue of the University of Oregon 241

454 correct estimate of the true value of β is 0.412.

36]

1000

$$|f_t, |f_t, \mathbb{V}$$
$$[40] \quad \langle \mathbf{N}(\mathbf{p}) \rangle = \langle \mathbf{N}(\mathbf{p}) \rangle_{\text{stat}} + \langle \mathbf{N}(\mathbf{p}) \rangle_{\text{dyn}}$$
$$\Lambda_{\infty}^{\infty} \quad \Lambda_{\infty}^{\infty} \{1, 1, 1, 1\};$$
$$m = 1 + \frac{1}{2} \frac{1}{\Gamma(1/2)} \frac{\Gamma(1/2)}{\Gamma(1/2)} \left(\frac{1}{2} \frac{1}{\Gamma(1/2)} \frac{\Gamma(1/2)}{\Gamma(1/2)} \right)$$

the DNA encoding mouse human diagnostic protein #27795.

[illegible]

ford supplement; medical imaging; diagnostic; qualitative; discrete; SS.

MEMORANDUM

W0200175067 A2 - 2V 7905/10070M
N2
N2
N2

1983 11 MAR 20 00:27:00.000000S 064921.7.
1983 11 MAR 20 00:27:00.000000S 064921.7.
1983 11 MAR 20 00:27:00.000000S 064921.7.

XXXXXX

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